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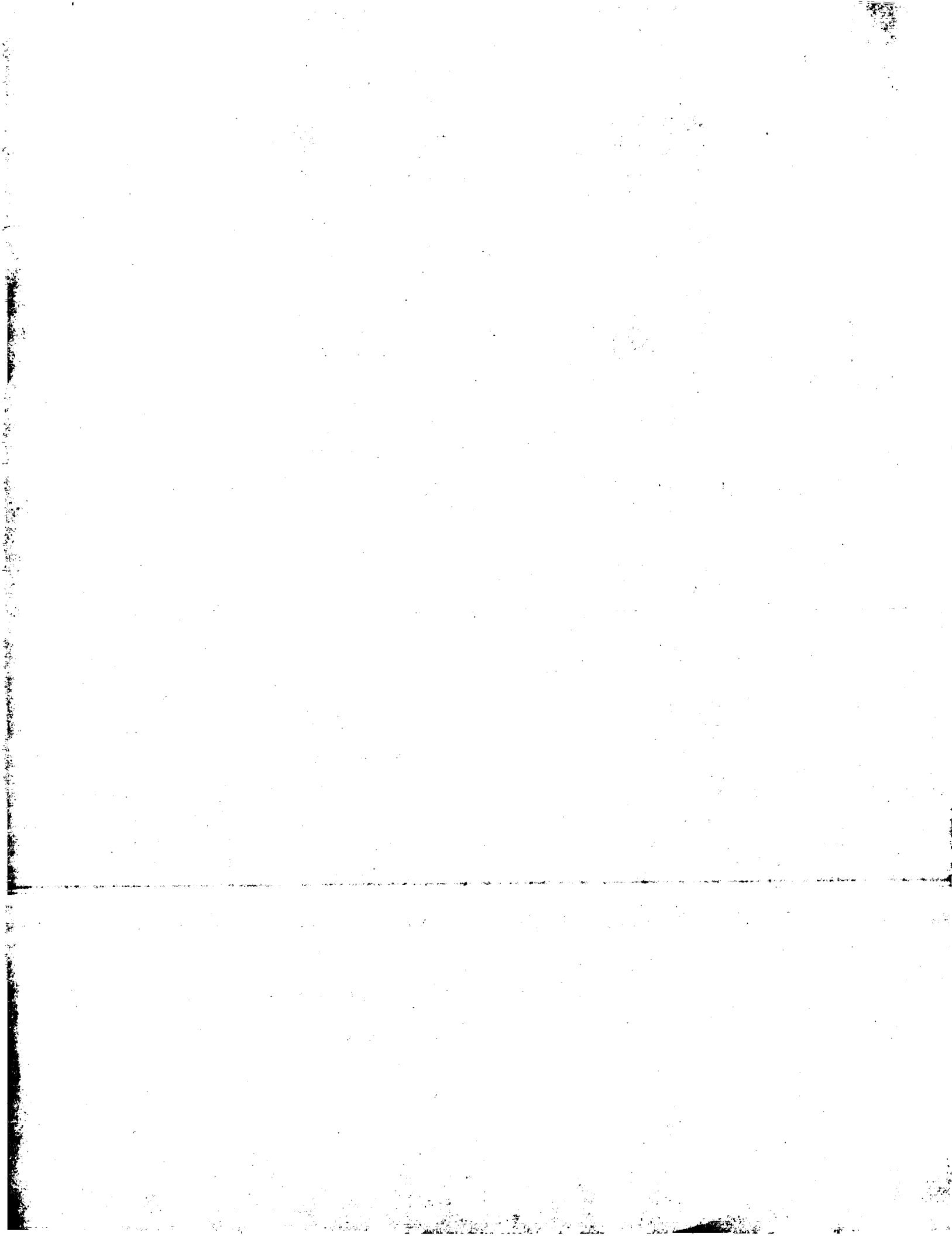
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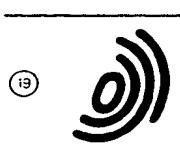
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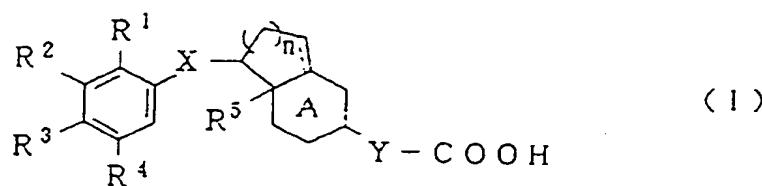
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(54) Carboxylic acid derivatives having 5 alpha-reductase inhibitory activity.

(57) Carboxylic acid derivatives of the formula:



are provided, which are useful for treatment of androgen dependent diseases such as benign prostatic hyperplasia, acne, seborrhea, female hirsutism, and male alopecia.

EP 0 579 223 A1

The present invention relates to novel carboxylic acid derivatives. More particularly, this invention relates to novel carboxylic acid derivatives having 5 α -reductase inhibitory activity useful for treating benign prostatic hyperplasia, acne, seborrhea, female hirsutism, prostatic carcinoma, male alopecia, or the like, caused by excessive production of dihydrotestosterone (hereinafter referred to as "DHT").

5 Androgen dependent diseases which represent unfavorable physiological symptoms such as benign prostatic hyperplasia, acne, seborrhea, female hirsutism, male alopecia, or the like are caused by excessive accumulation of androgenic hormones in metabolic system.

10 It has long been known that DHT is essential for differentiation, development and maintenance of prostatic tissue. It has also been well known that the active androgen of male target organs such as prostate 15 sebaceous gland, hair-root is DHT.

DHT is produced from testosterone by the action of a testosterone 5 α -hydrogenating enzyme, "5 α -reductase", in above target organs. Therefore, testosterone is a kind of pro-hormone in androgen-dependant tissues such as prostate gland, and 5 α -reductase plays an important role for the biosynthesis of DHT.

Importance of DHT concentration has recently been recognized in diseases which appear to be caused 20 by an excess of male hormone, and a lot of 5 α -reductase inhibitors have been reported, which include steroid derivatives such as 4-aza-steroid derivatives [JMC, 27, 1690 (1984)], 3-carboxylic acid steroid derivatives [Bioorganic chem, 17, 372 (1989)], JMC, 33, 937 (1990)], 3-phosphonic acid steroid derivatives [Japanese Patent Publication Kokai Hei 2-212499; Hei 2-225496], 3-sulfonic acid steroid derivatives [Japanese Patent Publication Kokai Hei 2-225494], 3-nitrosteroid derivatives [Japanese Patent Publication Kokai Hei 3-118325], non-steroid agents such as benzoylaminophenoxybutanoic acid derivatives [Japanese Patent Publication Kokai Hei 1-156950; Hei 1-139558], WS-9659 A and B originating from microorganisms [The Journal of Antibiotics, 1230, 1235, 1989], and the like.

The 5 α -reductase inhibitors listed above are classified in two groups, namely steroid derivatives and 25 non-steroid derivatives. The steroid derivatives have a problem of inducing side effects although they exhibit excellent pharmacological activities. On the other hand, non-steroid derivative having sufficient activity has not been discovered yet.

As the result of extensive study for providing non-steroid compounds having high activity as 5 α -reductase inhibitors, the present inventors have found that specific carboxylic acid derivatives have an excellent activity. The present invention has been completed on the basis of such finding.

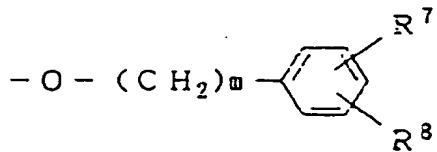
30 Thus, the present invention provides carboxylic acid derivatives of the following general formula (I):



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Wherein R¹, R², R³ and R⁴ each independently represent hydrogen atom, halogen atom, adamantyl group, optionally substituted C₁ - C₁₄ alkyl group, optionally substituted C₃ - C₁₀ cycloalkyl group, 45 optionally substituted C₁ - C₁₄ alkoxy group, optionally substituted heterocyclic group, - OR⁶ (R⁶ represents hydrogen atom, adamantyl group, optionally substituted C₃ - C₁₀ cycloalkyl group or optionally substituted heterocyclic group), or a group of the formula:

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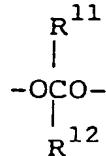


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wherein R⁷ and R⁸ each independently represent hydrogen atom, C₁ - C₆ alkyl group, C₃ - C₈ cycloalkyl group, - CONR⁹N¹⁰ (R⁹ and R¹⁰ each independently represent hydrogen atom or C₁ - C₆ alkyl group) or,

EP 0 579 223 A1

when R⁷ and R⁸ are adjacent, they may form C₁ - C₆ alkylene group, and m represents 0 or 1, or, the adjacent two substituents selected from R¹, R², R³ and R⁴ may form a group of the formula:



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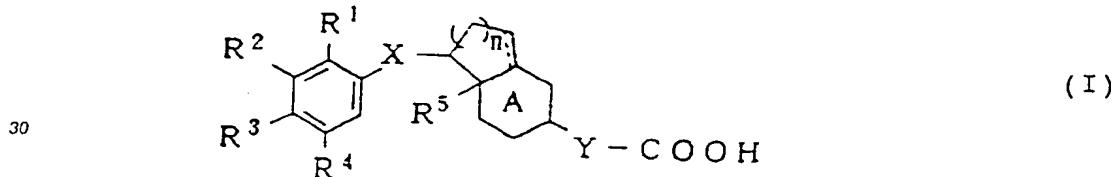
wherein R¹¹ and R¹² each independently represent hydrogen atom, C₁ - C₆ alkyl group or C₃ - C₈ cycloalkyl group or they may form, taken together, C₂ - C₈ alkylene group, a group of the formula: -OCH₂CH₂O- or optionally substituted C₃ - C₄ alkylene group,

15 R⁵ represents hydrogen atom or C₁ - C₅ alkyl group,
 X represents -CONR¹³- or -SO₂NR¹³- (R¹³ represents hydrogen atom or C₁ - C₆ alkyl group),
 Y represents a single bond, -OCH₂- or -CH=CH-,
 A ring may form benzene ring, cyclohexene ring or cyclohexadiene ring,
 the dotted line represents a single bond or double bond, and
 20 n represents 1 or 2, with the proviso that when the carbon atom to which R⁵ is attached has a double bond, then R⁵ is not present, or pharmaceutically acceptable salts thereof.

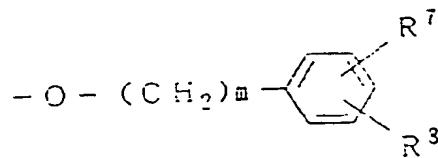
The present invention will be explained in detail below.

The present invention relates to the carboxylic acid derivatives of the following general formula (I):

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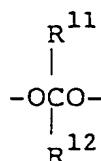


30 wherein R¹, R², R³ and R⁴ each independently represent hydrogen atom, halogen atom (iodine atom, fluorine atom, chlorine atom, bromine atom), adamantyl group, C₁ - C₁₄ alkyl group (methyl group, pentyl group, nonyl group, tetradecyl group, etc.) optionally having one or more substituents selected from 5 or 6 membered heterocyclic ring containing 1 or 2 hetero atoms selected from oxygen atom, sulfur atom and nitrogen atom, such as tetrahydrofuran ring, imidazoline ring, piperidine ring, dithian ring, thiomorpholine ring, etc., C₃ - C₁₀ cycloalkyl group (cyclopropyl group, cyclohexyl group, cyclodecyl group, etc.) and adamantyl group, C₃ - C₁₀ cycloalkyl group (cyclopropyl group, cyclohexyl group, cyclodecyl group, etc.) optionally having one or more substituents selected from C₁ - C₆ alkyl group (methyl group, butyl group, hexyl group, etc.), C₁ - C₁₄ alkoxy group (methoxy group, pentyloxy group, nonyloxy group, tetradecyloxy group, etc.) optionally having one or more substituents selected from 5 or 6 membered heterocyclic ring containing 1 or 2 hetero atoms selected from oxygen atom, sulfur atom and nitrogen atom, such as tetrahydrofuran ring, imidazoline ring, piperidine ring, dithian ring, thiomorpholine ring, etc., optionally having one or more substituents selected from C₁ - C₆ alkyl group (methyl group, butyl group, hexyl group, etc.), a group: -OR⁶ {R⁶ represents hydrogen atom, adamantyl group, C₃ - C₁₀ cycloalkyl group (cyclopropyl group, cyclohexyl group, cyclodecyl group, etc.) optionally having one or more substituents selected from C₁ - C₆ alkyl group (methyl group, butyl group, hexyl group, etc.) or 5 or 6 membered heterocyclic ring containing 1 or 2 hetero atoms selected from oxygen atom, sulfur atom and nitrogen atom (tetrahydrofuran ring, imidazoline ring, piperidine ring, dithian ring, thiomorpholine ring, etc.) 40 optionally having one or more substituents selected from C₁ - C₆ alkyl group (methyl group, butyl group, hexyl group, etc.) and C₂ - C₆ acyl group (acetyl group, isobutyryl group, isovaleryl group, etc.)} or a group:
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10 wherein R⁷ and R⁸ each independently represent hydrogen atom, C₁ - C₆ alkyl group (methyl group, butyl group, hexyl group, etc.), C₃ - C₈ cycloalkyl group (cyclopropyl group, cyclohexyl group, cyclooctyl group, etc.), -CONR⁹N¹⁰ [R⁹ and R¹⁰ each independently represent hydrogen atom or C₁ - C₆ alkyl group (methyl group, butyl group, hexyl group, etc.), or, when R⁷ and R⁸ are adjacent, they may form C₁ - C₆ alkylene group (methylene group, trimethylene group, hexamethylene group, etc.), and n represents 0 or 1}, or, the two adjacent substituents selected from R¹, R², R³ and R⁴ may form a group of the formula:

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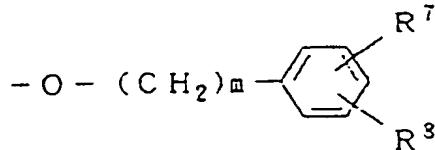
25 wherein R¹¹ and R¹² each independently represent hydrogen atom, C₁ - C₆ alkyl group (methyl group, butyl group, hexyl group, etc.) or C₃ - C₈ cycloalkyl group (cyclopropyl group, cyclopentyl group, cyclooctyl group, etc.) or, they may form C₂ - C₈ alkylene group (ethylene group, pentamethylene group, octamethylene group, etc.), a group: -OCH₂CH₂O- or C₃ - C₄ alkylene group (trimethylene group, tetramethylene group) optionally having one or more substituents selected from C₁ - C₆ alkyl group (methyl group, butyl group, hexyl group, etc.) and C₃ - C₈ cycloalkyl group (cyclopropyl group, cyclohexyl group, cyclooctyl group, etc.);

30 R⁵ represents hydrogen atom or C₁ - C₆ alkyl group (methyl group, butyl group, hexyl group, etc.), X represents -CONR¹³- or -SO₂NR¹³- [R¹³ represents hydrogen atom or C₁ - C₆ alkyl group (methyl group, butyl group, hexyl group, etc.)];

35 Y represents a single bond, -OCH₂- or -CH=CH-; A ring may form a benzene ring or cyclohexadiene ring; the dotted line represents a single bond or double bond; and n represents 1 or 2, with the proviso that, when the carbon to which R⁵ is attached has a double bond, then R⁵ is not present, or pharmaceutically acceptable salts thereof.

40 Of the compounds of the present invention, preferable is a compound of the formula (I) in which R¹, R², R³ and R⁴ each independently represent hydrogen atom; halogen atom; C₁ - C₁₄ alkyl group optionally having one or more substituents selected from 5 or 6 membered heterocyclic group containing 1 or 2 hetero atoms selected from oxygen atom, sulfur atom and nitrogen atom, and C₃ - C₁₀ cycloalkyl group; C₁ - C₁₄ alkoxy group optionally having one or more C₃ - C₁₀ cycloalkyl group; -OR⁶ [R⁶ represents C₃ - C₁₀ cycloalkyl group optionally having one or more C₁ - C₆ alkyl group; or 5 or 6 membered heterocyclic group containing 1 or 2 hetero atoms selected from oxygen atom, sulfur atom and nitrogen atom, and optionally having one or more C₁ - C₆ alkyl groups] or a group of the formula:

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EP 0 579 223 A1

wherein R⁷ and R⁸ each independently represent hydrogen atom, C₁ - C₆ alkyl group, C₃ - C₈ cycloalkyl group, - CONR⁹N¹⁰ (R⁹ and R¹⁰ each independently represent hydrogen atom or C₁ - C₆ alkyl group) or, when R⁷ and R⁸ are adjacent, they may form C₁ - C₆ alkylene group, and m represents 0 or 1), R⁵ represents C₁ - C₆ alkyl group, X represents -CONR¹³- or -SO₂NR¹³- (R¹³ represents hydrogen or C₁ - C₆ alkyl group), Y represents a single bond, -OCH₂-or -CH=CH-, A ring may form a benzene ring or cyclohexene ring, the dotted line represents a single bond or double bond, and n represents 1 or 2.

Particularly preferable is a compound of the formula (I) in which R¹, R², R³ and R⁴ each independently represent hydrogen atom; halogen atom; C₁ - C₁₄ alkyl group optionally having one or more C₃ - C₁₀ cycloalkyl groups; C₁ - C₁₄ alkoxy group optionally having one or more C₃ - C₁₀ cycloalkyl group; or -OR⁶ - (R⁶ represents C₃ - C₁₅ cycloalkyl group optionally having one or more C₁ - C₆ alkyl group); R⁵ represents C₁ - C₆ alkyl group, X represents -CONR¹³- (R¹³ represents hydrogen or C₁ - C₆ alkyl group); Y represents a single bond; A ring represents cyclohexene ring; the dotted line represents a single bond, and n represents 1 or 2.

Most preferable is a compound of the formula (I) in which R¹, R², R³ and R⁴ each independently represent hydrogen atom or C₁ - C₁₄ alkoxy group; R⁵ represents C₁ - C₆ alkyl group; X represents -CONR¹³- (R¹³ represents hydrogen atom); Y represents a single bond; A ring represents a cyclohexene ring; the dotted line represents a double bond; and n represents 2.

The compounds of the general formula (I) above may contain an asymmetric carbon atom, and racemic mixture and optical isomers are included in the present invention.

20 The compounds of the present invention are illustrated in Tables 1 - 4 below.

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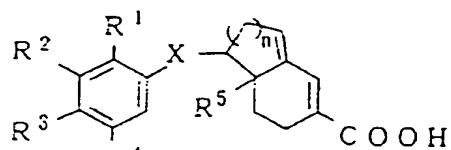
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Table 1

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
1	H	H		H	-Me	-CONH-	2
2	H	H		H	-Me	-CONH-	2
3	H	H		H	-Me	-CONH-	2
4	H	H		H	-Me	-CONH-	2
5	H	H		H	-Me	-CONH-	2
6	H	H		H	-Me	-CONMe-	2
7	H	H		H	-Me	-CONEt-	2
8	H	H		H	-Me	-CONH-	1
9	H	H		H	-Me	-CONMe-	1

Table 1 (continued)

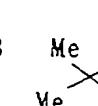
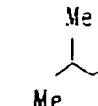
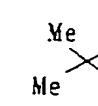
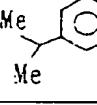
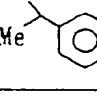
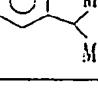
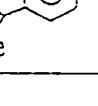
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
10	H		H	H	-Me	-CONH-	2
11	H		H	H	-Me	-CONH-	1
12			H	H	-Me	-CONH-	2
13	Me		H	H	-Me	-CONH-	2
14	Me		H	H	-Me	-CONH-	2
15	Me		H	H	-Me	-CONH-	1
16	H		H	H	-Me	-CONH-	2
17	H		H	H	-Me	-CONH-	2
18	H		H	H	-Me	-CONH-	2
19	H		H	H	-Me	-CONMe-	2

Table 1 (continued)

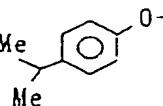
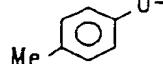
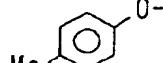
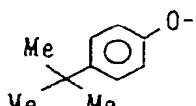
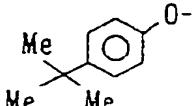
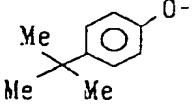
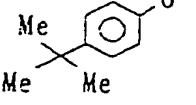
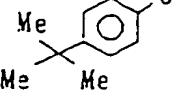
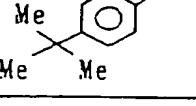
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
20	H		H	H	-Me	-CONH-	1
21	H		H	H	-Me	-CONH-	2
22	H		H	H	-Me	-CONH-	1
23	H		H	H	-Me	-CONMe-	2
24	H		H	H	-Me	-CONH-	2
25	H		H	H	-Me	-CONH-	1
26	H		H	H	-Me	-CONMe-	2
27	-OH		H	H	-Me	-CONH-	2
28	-Me		H	H	-Me	-CONH-	2
29	-Et		H	H	-Me	-CONH-	2

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
30	H		H	H	-Me	-CONH-	2
31	H		H	H	-Me	-CONH-	1
32	H		H	H	-Me	-CONMe-	2
33	H		H	H	-Me	-CONH-	2
34	H		H	H	-Me	-CONH-	1
35	H		H	H	-Me	-CONMe-	2
36	H		H	H	-Me	-CONH-	2
37	H		H	H	-Me	-CONH-	1
38	H		H	H	-Me	-CONH-	2
39	-Me		H	H	-Me	-CONH-	2

Table 1 (continued)

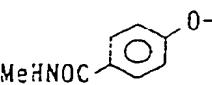
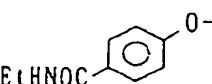
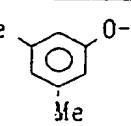
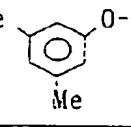
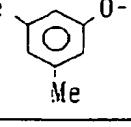
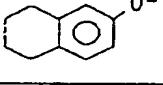
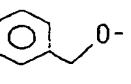
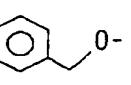
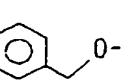
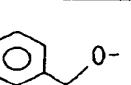
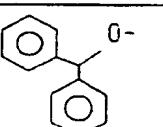
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
40	H		H	H	-Me	-CONH-	1
41	H		H	H	-Me	-CONH-	2
42	H		H	H	-Me	-CONH-	2
43	H		H	H	-Me	-CONH-	1
44	-OH		H	H	-Me	-CONH-	2
45	H		H	H	-Me	-CONH-	2
46	H		H	H	-Me	-CONH-	2
47	-Me		H	H	-Me	-CONH-	2
48	-OH		H	H	-Me	-CONH-	2
49	H		H	H	-Me	-CONH-	1
50	H		H	H	-Me	-CONH-	2

Table 1 (continued)

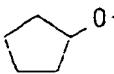
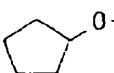
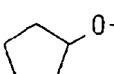
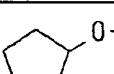
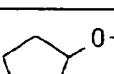
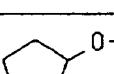
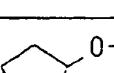
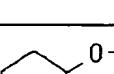
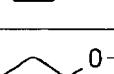
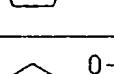
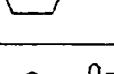
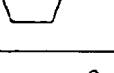
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
51	H		H	H	-Me	-CONH-	2
52	H		Me-	H	-Me	-CONH-	2
53	H		Et-	H	-Me	-CONH-	2
54	H		iPr-	H	-Me	-CONH-	2
55	H		Me-O-	H	-Me	-CONH-	2
56	H		Et-O-	H	-Me	-CONH-	2
57	H		iPr-	H	-Me	-CONH-	2
58	-OH		H	H	-Me	-CONH-	2
59	-F		H	H	-Me	-CONH-	2
60	-Cl		H	H	-Me	-CONH-	2
61	-Br		H	H	-Me	-CONH-	2
62	H		H	H	-Me	-CONH-	1

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
63	H		H	H	-Me	-CONH-	2
64	H		Me-	H	-Me	-CONH-	2
65	H		Et-	H	-Me	-CONH-	2
66	H		iPr-	H	-Me	-CONH-	2
67	H		Me-O-	H	-Me	-CONH-	2
68	H		Et-O-	H	-Me	-CONH-	2
69	H		iPr-O-	H	-Me	-CONH-	2
70	-OH		H	H	-Me	-CONH-	2
71	-Me		H	H	-Me	-CONH-	2
72	-F		H	H	-Me	-CONH-	2
73	-Cl		H	H	-Me	-CONH-	2
74	-Br		H	H	-Me	-CONH-	2

Table 1 (continued)

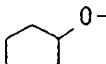
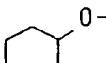
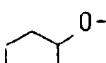
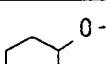
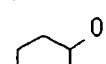
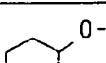
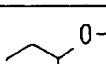
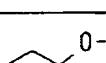
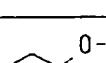
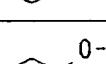
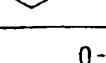
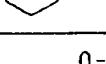
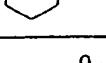
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
75	H		H	F-	-Me	-CONH-	2
76	H		H	Cl-	-Me	-CONH-	2
77	H		H	Br-	-Me	-CONH-	2
78	H		H		-Me	-CONH-	2
79	H		H	H	-Me	-CONMe-	2
80	H		Me-	H	-Me	-CONMe-	2
81	H		Et-	H	-Me	-CONMe-	2
82	H		iPr-	H	-Me	-CONMe-	2
83	H		Me-O-	H	-Me	-CONMe-	2
84	H		Et-O-	H	-Me	-CONMe-	2
85	H		iPr-O-	H	-Me	-CONMe-	2
86	-OH		H	H	-Me	-CONMe-	2

Table 1 (continued)

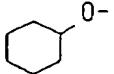
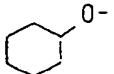
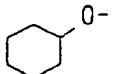
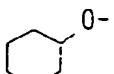
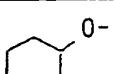
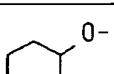
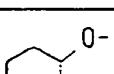
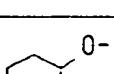
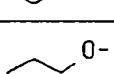
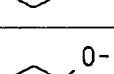
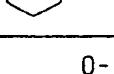
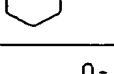
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
87	H		H	H	-Me	-CONH-	1
88	H		Me-	H	-Me	-CONH-	1
89	H		Et-	H	-Me	-CONH-	1
90	H		iPr-	H	-Me	-CONH-	1
91	H		Me-O-	H	-Me	-CONH-	1
92	H		Et-O-	H	-Me	-CONH-	1
93	H		iPr-O-	H	-Me	-CONH-	1
94	-OH		H	H	-Me	-CONH-	1
95	-Me		H	H	-Me	-CONH-	1
96	-F		H	H	-Me	-CONH-	1
97	-Cl		H	H	-Me	-CONH-	1
98	-Br		H	H	-Me	-CONH-	1

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
99	H		H	F-	-Me	-CONH-	1
100	H		H	Cl-	-Me	-CONH-	1
101	H		H	Br-	-Me	-CONH-	1
102	H		H		-Me	-CONH-	1
103	H		H	H	-Me	-CONMe-	1
104	H		Me-	H	-Me	-CONMe-	1
105	H		Et-	H	-Me	-CONMe-	1
106	H		iPr-	H	-Me	-CONMe-	1
107	H		Me-O-	H	-Me	-CONMe-	1
108	H		Et-O-	H	-Me	-CONMe-	1
109	H		iPr-C-	H	-Me	-CONMe-	1
110	-OH		H	H	-Me	-CONMe-	1

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
5 111	H		H	H	-Me	-CONH-	2
10 112	H		Me-	H	-Me	-CONH-	2
15 113	H		Et-	H	-Me	-CONH-	2
20 114	H		iPr-	H	-Me	-CONH-	2
25 115	H		Me-O-	H	-Me	-CONH-	2
30 116	H		Et-O-	H	-Me	-CONH-	2
35 117	H		iPr-O-	H	-Me	-CONH-	2
40 118	-OH		H	H	-Me	-CONH-	2
45 119	-Me		H	H	-Me	-CONH-	2
50 120	-F		H	H	-Me	-CONH-	2
55 121	-Cl		H	H	-Me	-CONH-	2
60 122	-Br		H	H	-Me	-CONH-	2

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
123	H		H	F-	-Me	-CONH-	2
124	H		H	Cl-	-Me	-CONH-	2
125	H		H	Br-	-Me	-CONH-	2
126	H		H		-Me	-CONH-	2
127	H		H	H	-Me	-CONMe-	2
128	H		Me-	H	-Me	-CONMe-	2
129	H		Et-	H	-Me	-CONMe-	2
130	H		iPr-	H	-Me	-CONMe-	2
131	H		Me-O-	H	-Me	-CONMe-	2
132	H		Et-O-	H	-Me	-CONMe-	2
133	H		iPr-O-	H	-Me	-CONMe-	2
134	-OH		H	H	-Me	-CONMe-	2

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
135	H		H	H	-Me	-CONH-	1
136	H		Me-	H	-Me	-CONH-	1
137	H		Et-	H	-Me	-CONH-	1
138	H		iPr-	H	-Me	-CONH-	1
139	H		Me-O-	H	-Me	-CONH-	1
140	H		Et-O-	H	-Me	-CONH-	1
141	H		iPr-O-	H	-Me	-CONH-	1
142	-OH		H	H	-Me	-CONH-	1
143	-Me		H	H	-Me	-CONH-	1
144	-F		H	H	-Me	-CONH-	1
145	-Cl		H	H	-Me	-CONH-	1
146	-Br		H	H	-Me	-CONH-	1

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
5 147	H		H	F-	-Me	-CONH-	1
10 148	H		H	Cl-	-Me	-CONH-	1
15 149	H		H	Br-	-Me	-CONH-	1
20 150	H		H		-Me	-CONH-	1
25 151	H		H	H	-Me	-CONMe-	1
30 152	H		Me-	H	-Me	-CONMe-	1
35 153	H		Et-	H	-Me	-CONMe-	1
40 154	H		iPr-	H	-Me	-CONMe-	1
45 155	H		Me-O-	H	-Me	-CONMe-	1
50 156	H		Et-O-	H	-Me	-CONMe-	1
55 157	H		iPr-O-	H	-Me	-CONMe-	1
56 158	-OH		H	H	-Me	-CONMe-	1

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
159	H		H	H	-Me	-CONH-	2
160	H		Me-	H	-Me	-CONH-	2
161	H		Et-	H	-Me	-CONH-	2
162	H		iPr-	H	-Me	-CONH-	2
163	H		Me-O-	H	-Me	-CONH-	2
164	H		Et-O-	H	-Me	-CONH-	2
165	H		iPr-O-	H	-Me	-CONH-	2
166	-OH		H	H	-Me	-CONH-	2
167	-Me		H	H	-Me	-CONH-	2
168	-F		H	H	-Me	-CONH-	2
169	-Cl		H	H	-Me	-CONH-	2
170	-Br		H	H	-Me	-CONH-	2

Table 1 (continued)

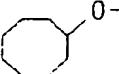
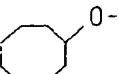
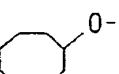
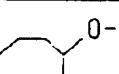
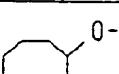
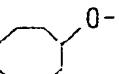
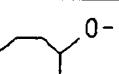
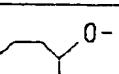
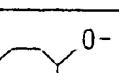
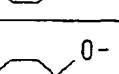
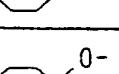
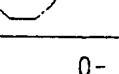
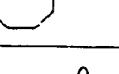
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
171	H		H	F-	-Me	-CONH-	2
172	H		H	Cl-	-Me	-CONH-	2
173	H		H	Br-	-Me	-CONH-	2
174	H		H		-Me	-CONH-	2
175	H		H	H	-Me	-CONMe-	2
176	H		Me-	H	-Me	-CONMe-	2
177	H		Et-	H	-Me	-CONMe-	2
178	H		iPr-	H	-Me	-CONMe-	2
179	H		Me-O-	H	-Me	-CONMe-	2
180	H		Et-O-	H	-Me	-CONMe-	2
181	H		iPr-O-	H	-Me	-CONMe-	2
182	-OH		H	H	-Me	-CONMe-	2

Table 1 (continued)

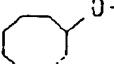
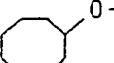
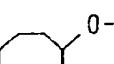
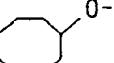
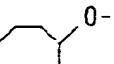
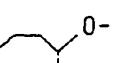
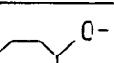
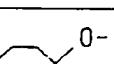
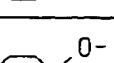
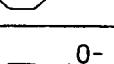
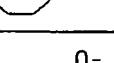
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
183	H			H	H	-Me	-CONH-
184	H			Me-	H	-Me	-CONH-
185	H			Et-	H	-Me	-CONH-
186	H			iPr-	H	-Me	-CONH-
187	H			Me-O-	H	-Me	-CONH-
188	H			Et-O-	H	-Me	-CONH-
189	H			iPr-O-	H	-Me	-CONH-
190	-OH			H	H	-Me	-CONH-
191	-Me			H	H	-Me	-CONH-
192	-F			H	H	-Me	-CONH-
193	-Cl			H	H	-Me	-CONH-
194	-Br			H	H	-Me	-CONH-

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
195	H		H	F-	-Me	-CONH-	1
196	H		H	Cl-	-Me	-CONH-	1
197	H		H	Br-	-Me	-CONH-	1
198	H		H		-Me	-CONH-	1
199	H		H	H	-Me	-CONMe-	1
200	H		Me-	H	-Me	-CONMe-	1
201	H		Et-	H	-Me	-CONMe-	1
202	H		iPr-	H	-Me	-CONMe-	1
203	H		Me-O-	H	-Me	-CONMe-	1
204	H		Et-O-	H	-Me	-CONMe-	1
205	H		iPr-O-	H	-Me	-CONMe-	1
206	-OH		H	H	-Me	-CONMe-	1

Table 1 (continued)

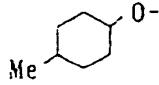
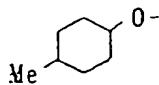
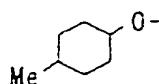
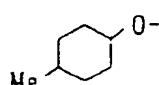
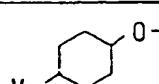
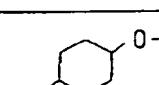
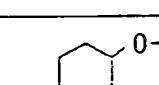
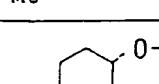
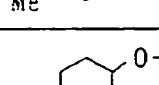
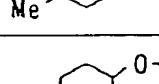
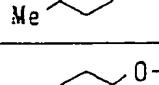
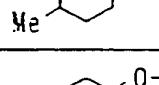
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
207	H		H	H	-Me	-CONH-	2
208	H		Me-	H	-Me	-CONH-	2
209	H		Et-	H	-Me	-CONH-	2
210	H		iPr-	H	-Me	-CONH-	2
211	H		Me-O-	H	-Me	-CONH-	2
212	H		Et-O-	H	-Me	-CONH-	2
213	H		iPr-O-	H	-Me	-CONH-	2
214	-OH		H	H	-Me	-CONH-	2
215	-Me		H	H	-Me	-CONH-	2
216	-F		H	H	-Me	-CONH-	2
217	-Cl		H	H	-Me	-CONH-	2
218	-Br		H	H	-Me	-CONH-	2

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
219	H		H	F-	-Me	-CONH-	2
220	H		H	Cl-	-Me	-CONH-	2
221	H		H	Br-	-Me	-CONH-	2
222	H		H		-Me	-CONH-	2
223	H		H	H	-Me	-CONMe-	2
224	H		Me-	H	-Me	-CONMe-	2
225	H		Et-	E	-Me	-CONMe-	2
226	H		iPr-	H	-Me	-CONMe-	2
227	H		Me-O-	H	-Me	-CONMe-	2
228	H		Et-O-	H	-Me	-CONMe-	2
229	H		iPr-O-	H	-Me	-CONMe-	2
230	-OH		H	H	-Me	-CONMe-	2

Table 1 (continued)

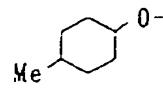
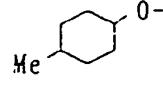
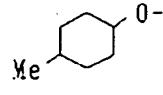
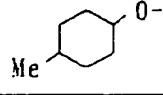
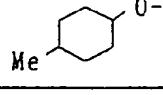
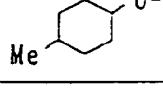
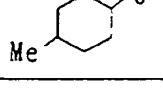
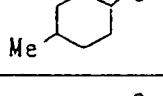
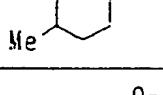
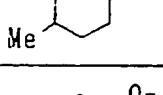
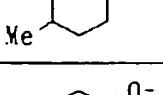
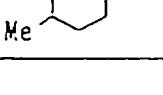
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
231	H		H	H	-Me	-CONH-	1
232	H		Me-	H	-Me	-CONH-	1
233	H		Et-	H	-Me	-CONH-	1
234	H		iPr-	H	-Me	-CONH-	1
235	H		Me-O-	H	-Me	-CONH-	1
236	H		Et-O-	H	-Me	-CONH-	1
237	H		iPr-O-	H	-Me	-CONH-	1
238	-OH		H	H	-Me	-CONH-	1
239	-Me		H	H	-Me	-CONH-	1
240	-F		H	H	-Me	-CONH-	1
241	-Cl		H	H	-Me	-CONH-	1
242	-Br		H	H	-Me	-CONH-	1

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
243	H		H	F-	-Me	-CONH-	1
244	H		H	Cl-	-Me	-CONH-	1
245	H		H	Br-	-Me	-CONH-	1
246	H		H		-Me	-CONH-	1
247	H		H	H	-Me	-CONMe-	1
248	H		Me-	H	-Me	-CONMe-	1
249	H		Et-	H	-Me	-CONMe-	1
250	H		iPr-	H	-Me	-CONMe-	1
251	H		Me-O-	H	-Me	-CONMe-	1
252	H		Et-O-	H	-Me	-CONMe-	1
253	H		iPr-O-	H	-Me	-CONMe-	1
254	-OH		H	H	-Me	-CONMe-	1

Table 1 (continued)

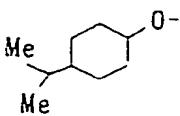
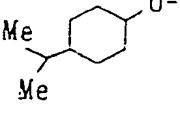
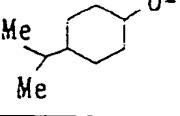
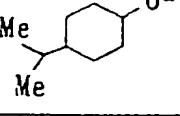
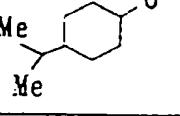
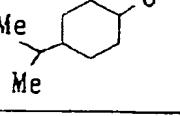
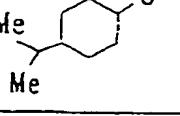
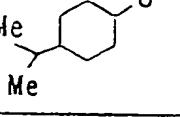
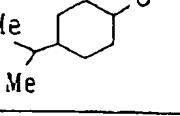
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
255	H		H	H	-Me	-CONH-	2
256	H		Me-	H	-Me	-CONH-	2
257	H		Et-	H	-Me	-CONH-	2
258	H		iPr-	H	-Me	-CONH-	2
259	H		Me-O-	H	-Me	-CONH-	2
260	H		Et-O-	H	-Me	-CONH-	2
261	H		iPr-O-	H	-Me	-CONH-	2
262	-OH		H	H	-Me	-CONH-	2
263	-Me		H	H	-Me	-CONH-	2

Table 1 (continued)

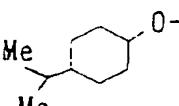
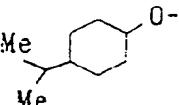
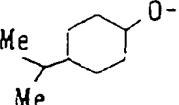
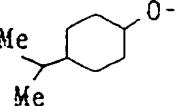
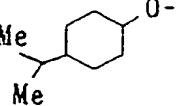
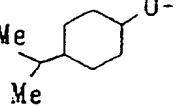
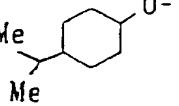
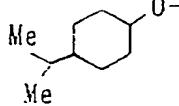
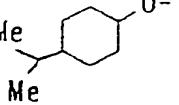
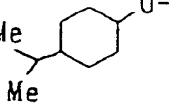
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
264	-F		H	H	-Me	-CONH-	2
265	-Cl		H	H	-Me	-CONH-	2
266	-Br		H	H	-Me	-CONH-	2
267	H		H	F-	-Me	-CONH-	2
268	H		H	Cl-	-Me	-CONH-	2
269	H		H	Br-	-Me	-CONH-	2
270	H			-Me	-CONH-	2	
271	H		H	H	-Me	-CONMe-	2
272	H		Me-	H	-Me	-CONMe-	2

Table 1 (continued)

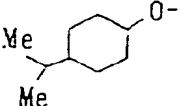
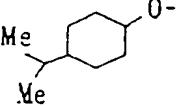
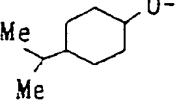
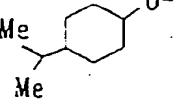
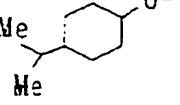
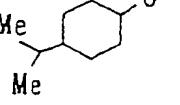
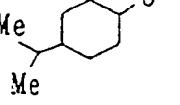
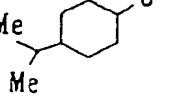
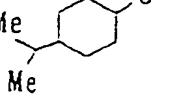
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
273	H		Et-	H	-Me	-CONMe-	2
274	H		iPr-	H	-Me	-CONMe-	2
275	H		Me-O-	H	-Me	-CONMe-	2
276	H		Et-O-	H	-Me	-CONMe-	2
277	H		iPr-O-	H	-Me	-CONMe-	2
278	-OH		H	H	-Me	-CONMe-	2
279	H		H	H	-Me	-CONH-	1
280	H		Me-	H	-Me	-CONH-	1
281	H		Et-	H	-Me	-CONH-	1

Table 1 (continued)

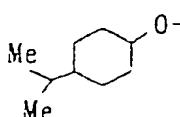
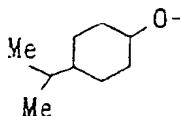
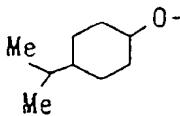
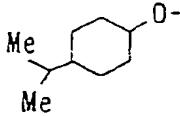
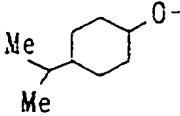
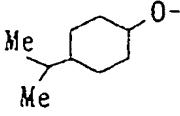
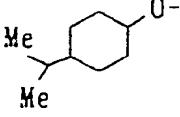
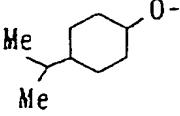
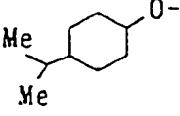
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
282	H		iPr-	H	-Me	-CONH-	1
283	H		Me-O-	H	-Me	-CONH-	1
284	H		Et-O-	H	-Me	-CONH-	1
285	H		iPr-O-	H	-Me	-CONH-	1
286	-OH		H	H	-Me	-CONH-	1
287	-Me		H	H	-Me	-CONH-	1
288	-F		H	H	-Me	-CONH-	1
289	-Cl		H	H	-Me	-CONH-	1
290	-Br		H	H	-Me	-CONH-	1

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
291	H		H	F-	-Me	-CONH-	1
292	H		H	Cl-	-Me	-CONH-	1
293	H		H	Br-	-Me	-CONH-	1
294	H		H		-Me	-CONH-	1
295	H		H	H	-Me	-CONMe-	1
296	H		Me-	H	-Me	-CONMe-	1
297	H		Et-	H	-Me	-CONMe-	1
298	H		iPr-	H	-Me	-CONMe-	1
299	H		Me-O-	H	-Me	-CONMe-	1

Table 1 (continued)

5 Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
10 300	H		Et-O-	H	-Me	-CONMe-	1
15 301	H		iPr-O-	H	-Me	-CONMe-	1
20 302	-OH		H	H	-Me	-CONMe-	1
25 303	H		H	H	-Me	-CONH-	2
30 304	H		Me-	H	-Me	-CONH-	2
35 305	H		Et-	H	-Me	-CONH-	2
40 306	H		iPr-	H	-Me	-CONH-	2
45 307	H		Me-O-	H	-Me	-CONH-	2
50 308	H		Bt-O-	H	-Me	-CONH-	2

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n	
309	H	Me Me		iPr-O-	H	-Me	-CONH-	2
310	-OH	Me Me		H	H	-Me	-CONH-	2
311	-Me	Me Me		H	H	-Me	-CONH-	2
312	-F	Me Me		H	H	-Me	-CONH-	2
313	-Cl	Me Me		H	H	-Me	-CONH-	2
314	-Br	Me Me		H	H	-Me	-CONH-	2
315	H	Me Me		H	F-	-Me	-CONH-	2
316	H	Me Me		H	Cl-	-Me	-CONH-	2
317	H	Me Me		H	Br-	-Me	-CONH-	2

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
318	H				-Me	-CONH-	2
319	H		H	H	-Me	-CONMe-	2
320	H		Et-	H	-Me	-CONMe-	2
321	H		iPr-	H	-Me	-CONMe-	2
322	H		Et-O-	H	-Me	-CONMe-	2
323	H		Me-O-	H	-Me	-CONMe-	2
324	H		Et-O-	H	-Me	-CONMe-	2
325	H		iPr-O-	H	-Me	-CONMe-	2
326	-OH		H	H	-Me	-CONMe-	2

Table 1 (continued)

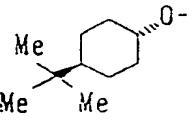
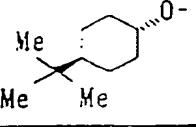
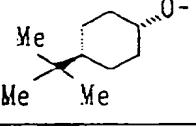
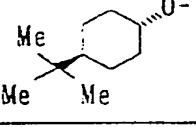
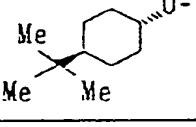
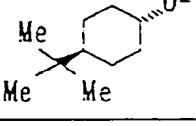
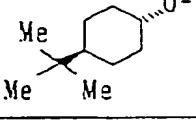
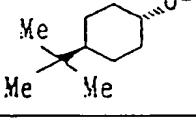
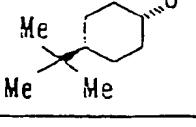
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
327	H	Me 	H	H	-Me	-CONH-	1
328	H	Me 	Me-	H	-Me	-CONH-	1
329	H	Me 	Et-	H	-Me	-CONH-	1
330	H	Me 	iPr-	H	-Me	-CONH-	1
331	H	Me 	Me-O-	H	-Me	-CONH-	1
332	H	Me 	Et-O-	H	-Me	-CONH-	1
333	H	Me 	iPr-O-	H	-Me	-CONH-	1
334	-OH	Me 	H	H	-Me	-CONH-	1
335	-Me	Me 	H	H	-Me	-CONH-	1

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
336	-F		H	H	-Me	-CONH-	1
337	-Cl		H	H	-Me	-CONH-	1
338	-Br		H	H	-Me	-CONH-	1
339	H		H	F-	-Me	-CONH-	1
340	H		H	Cl-	-Me	-CONH-	1
341	H		H	Br-	-Me	-CONH-	1
342	H		H		-Me	-CONH-	1
343	H		H	H	-Me	-CONMe-	1
344	H		Me-	H	-Me	-CONMe-	1

Table 1 (continued)

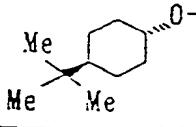
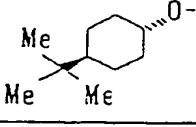
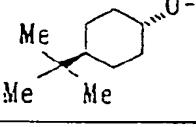
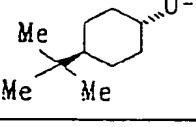
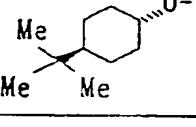
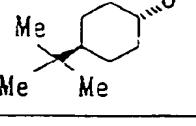
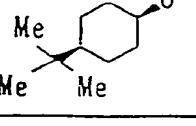
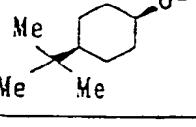
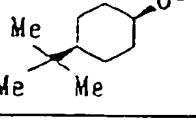
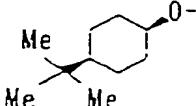
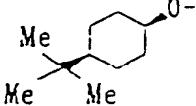
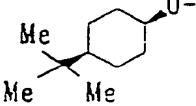
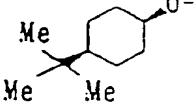
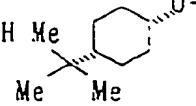
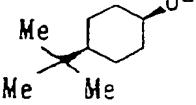
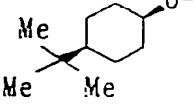
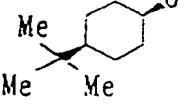
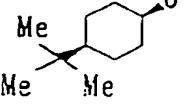
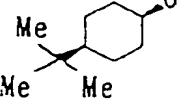
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
345	H	Me Me		Et-	H	-Me	-CONMe-
346	H	Me Me		iPr-	H	-Me	-CONMe-
347	H	Me Me		Me-O-	H	-Me	-CONMe-
348	H	Me Me		Et-O-	H	-Me	-CONMe-
349	H	Me Me		iPr-O-	H	-Me	-CONMe-
350	-OH	Me Me		H	H	-Me	-CONMe-
351	H	Me Me		H	H	-Me	-CONH-
352	H	Me Me		Me-	H	-Me	-CONH-
353	H	Me Me		Et-	H	-Me	-CONH-

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
354	H		iPr-	H	-Me	-CONH-	2
355	H		Me-O-	H	-Me	-CONH-	2
356	H		Et-O-	H	-Me	-CONH-	2
357	H		iPr-O-	H	-Me	-CONH-	2
358	-OH		H	H	-Me	-CONH-	2
359	-Me		H	H	-Me	-CONH-	2
360	-F		H	H	-Me	-CONH-	2
361	-Cl		H	H	-Me	-CONH-	2
362	-Br		H	H	-Me	-CONH-	2

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
363	H		H	F-	-Me	-CONH-	2
364	H		H	Cl-	-Me	-CONH-	2
365	H		H	Br-	-Me	-CONH-	2
366	H		H		-Me	-CONH-	2
367	H		H	H	-Me	-CONMe-	2
368	H		Me-	H	-Me	-CONMe-	2
369	H		Et-	H	-Me	-CONMe-	2
370	H		iPr-	H	-Me	-CONMe-	2
371	H		Me-O-	H	-Me	-CONMe-	2

Compd.
No.Table 1 (continued)

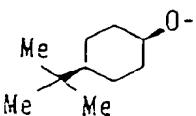
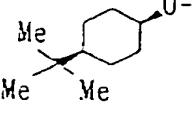
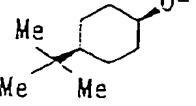
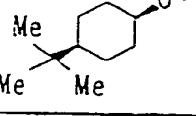
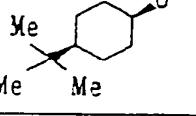
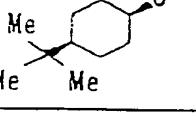
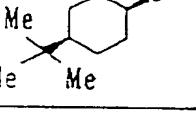
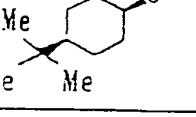
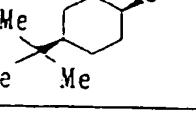
	R ¹	R ²	R ³	R ⁴	R ⁵	X	n	
5	372	H		Et-O-	H	-Me	-CONMe-	2
10	373	H		iPr-O-	H	-Me	-CONMe-	2
15	374	-OH		H	H	-Me	-CONMe-	2
20	375	H		H	H	-Me	-CONH-	1
25	376	H		Me-	H	-Me	-CONH-	1
30	377	H		Et-	H	-Me	-CONH-	1
35	378	H		iPr-	H	-Me	-CONH-	1
40	379	H		Me-O-	H	-Me	-CONH-	1
45	380	H		Et-O-	H	-Me	-CONH-	1
50								

Table 1 (continued)

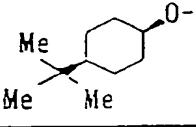
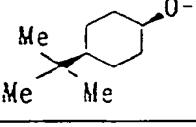
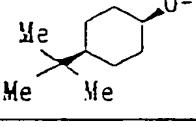
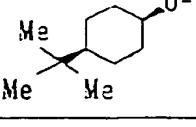
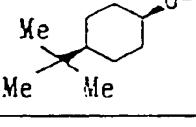
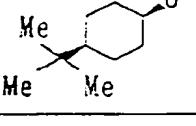
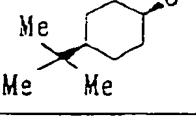
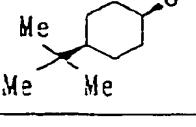
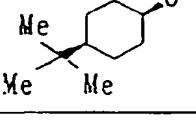
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
381	H	Me 	iPr-O-	H	-Me	-CONH-	1
382	-OH	Me 	H	H	-Me	-CONH-	1
383	-Me	Me 	H	H	-Me	-CCNH-	1
384	-F	Me 	H	H	-Me	-CONH-	1
385	-Cl	Me 	H	H	-Me	-CONH-	1
386	-Br	Me 	H	H	-Me	-CONH-	1
387	H	Me 	H	F-	-Me	-CONH-	1
388	H	Me 	H	Cl-	-Me	-CONH-	1
389	H	Me 	H	Br-	-Me	-CONH-	1

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
390	H				-Me	-CONH-	1
391	H		H	H	-Me	-CONMe-	1
392	H		Me-	H	-Me	-CONMe-	1
393	H		Et-	H	-Me	-CONMe-	1
394	H		iPr-	H	-Me	-CONMe-	1
395	H		Me-O-	H	-Me	-CONMe-	1
396	H		Et-O-	H	-Me	-CONMe-	1
397	H		iPr-O-	H	-Me	-CONMe-	1
398	-OH			H	-Me	-CONMe-	1
399	H			H	-Me	-CONH-	2

Table 1 (continued)

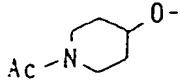
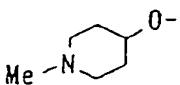
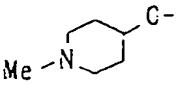
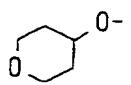
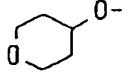
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
400	H		H	H	-Me	-CONH-	1
401	H		H	H	-Me	-CONH-	2
402	H		H	H	-Me	-CONH-	1
403	H		H	H	-Me	-CONH-	2
404	H		H	H	-Me	-CONH-	1
405	H	Me-	H	H	-Me	-CONH-	2
406	H	Et-	H	H	-Me	-CONH-	2
407	H	nPr-	H	H	-Me	-CONH-	2
408	H	iPr-	H	H	-Me	-CONH-	2
409	H	Me-O-	H	H	-Me	-CONH-	2
410	H	Et-O-	H	H	-Me	-CONH-	2
411	H	nPr-O-	H	H	-Me	-CONH-	2

Table 1 (continued)

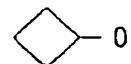
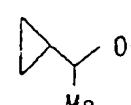
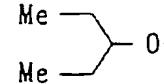
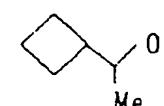
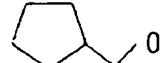
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
5 412	H	iPr-O-	H	H	-Me	-CONH-	2
10 413	H	 O-	H	H	-Me	-CONH-	2
15 414	H	nBu-O-	H	H	-Me	-CONH-	2
20 415	H	iBu-O-	H	H	-Me	-CONH-	2
25 416	H	 O-	H	H	-Me	-CONH-	2
30 417	H	 O-	H	H	-Me	-CONH-	2
35 418	H	 O-	H	H	-Me	-CONH-	2
40 419	H	 O-	H	H	-Me	-CONH-	2
45 420	H	 O-	H	H	-Me	-CONH-	2
50 421	H	 O-	H	H	-Me	-CONH-	2
55 422	H	 O-	H	H	-Me	-CONH-	2

Table 1 (continued)

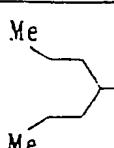
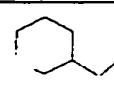
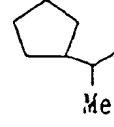
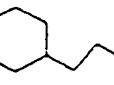
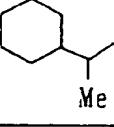
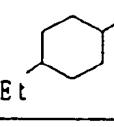
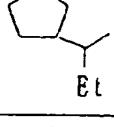
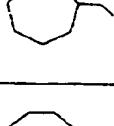
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
423	H		O-	H	H	-Me	-CONH-
424	H		O-	H	H	-Me	-CONH-
425	H		O-	H	H	-Me	-CONH-
426	H		O-	H	H	-Me	-CONH-
427	H		O-	H	H	-Me	-CONH-
428	H		O-	H	H	-Me	-CONH-
429	H		O-	H	H	-Me	-CONH-
430	H		O-	H	H	-Me	-CONH-
431	H		O-	H	H	-Me	-CONH-
432	H		O-	H	H	-Me	-CONH-

Table 1 (continued)

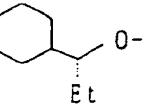
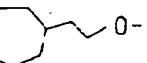
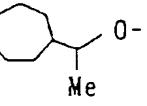
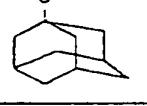
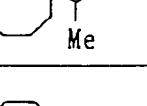
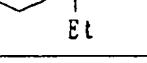
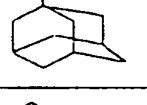
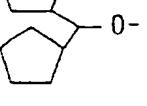
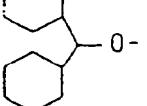
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
433	H		O-	H	H	-Me	-CONH-
434	H		O-	H	H	-Me	-CONH-
435	H		O-	H	H	-Me	-CONH-
436	H		O-	H	H	-Me	-CONH-
437	H		O-	H	H	-Me	-CONH-
438	H		O-	H	H	-Me	-CONH-
439	H		O-	H	H	-Me	-CONH-
440	H		O-	H	H	-Me	-CONH-
441	H		O-	H	H	-Me	-CONH-
442	H		O-	H	H	-Me	-CONH-

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
443	H	Me-	H	H	-Me	-CONH-	1
444	H	Et-	H	H	-Me	-CONH-	1
445	H	nPr-	H	H	-Me	-CONH-	1
446	H	iPr-	H	H	-Me	-CONH-	1
447	H	Me-O-	H	H	-Me	-CONH-	1
448	H	Et-O-	H	H	-Me	-CONH-	1
449	H	nPr-O-	H	H	-Me	-CONH-	1
450	H	iPr-O-	H	H	-Me	-CONH-	1
451	H	 O-	H	H	-Me	-CONH-	1
452	H	nBu-O-	H	H	-Me	-CONH-	1
453	H	iBu-O-	H	H	-Me	-CONH-	1
454	H	 O-	H	H	-Me	-CONH-	1

Table 1 (continued)

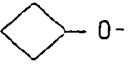
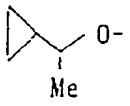
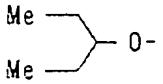
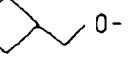
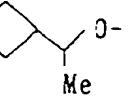
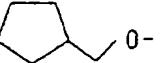
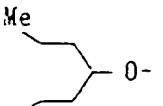
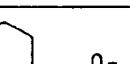
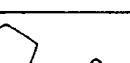
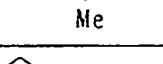
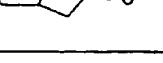
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
455	H		O-	H	H	-Me	-CONH-
456	H		O-	H	H	-Me	-CONH-
457	H		O-	H	H	-Me	-CONH-
458	H		O-	H	H	-Me	-CONH-
459	H		O-	H	H	-Me	-CONH-
460	H		O-	H	H	-Me	-CONH-
461	H		O-	H	H	-Me	-CONH-
462	H		O-	H	H	-Me	-CONH-
463	H		O-	H	H	-Me	-CONH-
464	H		O-	H	H	-Me	-CONH-
465	H		O-	H	H	-Me	-CONH-

Table 1 (continued)

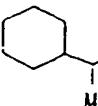
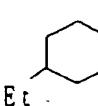
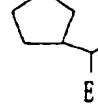
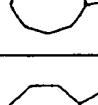
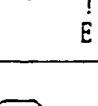
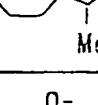
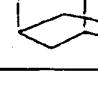
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
466	H		O-	H	H	-Me	-CONH-
467	H		O-	H	H	-Me	-CONH-
468	H		O-	H	H	-Me	-CONH-
469	H		O-	H	H	-Me	-CONH-
470	H		O-	H	H	-Me	-CONH-
471	H		O-	H	H	-Me	-CONH-
472	H		O-	H	H	-Me	-CONH-
473	H		O-	H	H	-Me	-CONH-
474	H		O-	H	H	-Me	-CONH-
475	H		O-	H	H	-Me	-CONH-

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
476	H		H	H	-Me	-CONH-	1
477	H		H	H	-Me	-CONH-	1
478	H		H	H	-Me	-CONH-	1
479	H		H	H	-Me	-CONH-	1
480	H		H	H	-Me	-CONH-	1
481	H		H	H	-Me	-CONH-	2
482	H		Me-	H	-Me	-CONH-	2
483	H		Et-	H	-Me	-CONH-	2
484	H		iPr-	H	-Me	-CONH-	2

Table 1 (continued)

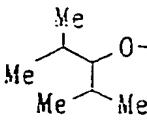
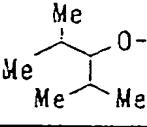
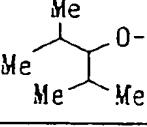
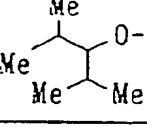
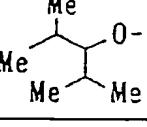
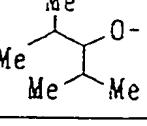
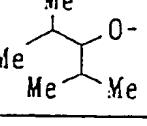
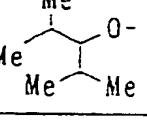
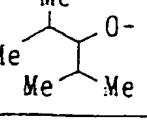
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n	
485	H		Me-O-	Me-O-	H	-Me	-CONH-	2
486	H		Et-O-	H	-Me	-CONH-	2	
487	H		iPr-O-	H	-Me	-CONH-	2	
488	-OH		H	H	-Me	-CONH-	2	
489	-Me		H	H	-Me	-CONH-	2	
490	-F		H	H	-Me	-CONH-	2	
491	-Cl		H	H	-Me	-CONH-	2	
492	-Br		H	H	-Me	-CONH-	2	
493	H		H	F-	-Me	-CONH-	2	

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
494	H		H	Cl-	-Me	-CONH-	2
495	H		H	Br-	-Me	-CONH-	2
496	H		H		-Me	-CONH-	2
497	H		H	H	-Me	-CONMe-	2
498	H		Me-	H	-Me	-CONMe-	2
499	H		Et-	H	-Me	-CONMe-	2
500	H		iPr-	H	-Me	-CONMe-	2
501	H		Me-O-	H	-Me	-CONMe-	2
502	H		Et-O-	H	-Me	-CONMe-	2

Table 1 (continued)

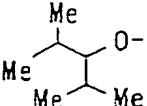
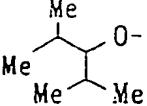
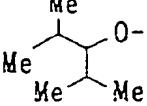
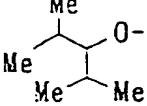
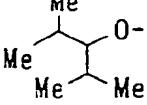
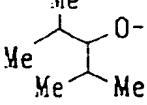
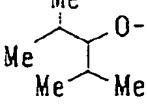
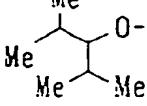
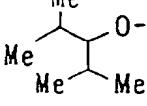
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
503	H		iPr-O-	H	-Me	-CONMe-	2
504	-OH		H	H	-Me	-CONMe-	2
505	H		H	H	-Me	-CONH-	1
506	H		Me-	H	-Me	-CONH-	1
507	H		Et-	H	-Me	-CONH-	1
508	H		iPr-	H	-Me	-CONH-	1
509	H		Me-O-	H	-Me	-CONH-	1
510	H		Et-O-	H	-Me	-CONH-	1
511	H		iPr-O-	H	-Me	-CONH-	1

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
512	-OH		H	H	-Me	-CONH-	1
513	-Me		H	H	-Me	-CONH-	1
514	-F		H	H	-Me	-CONH-	1
515	-Cl		H	H	-Me	-CONH-	1
516	-Br		H	H	-Me	-CONH-	1
517	H		H	F-	-Me	-CONH-	1
518	H		H	Cl-	-Me	-CONH-	1
519	H		H	Br-	-Me	-CONH-	1
520	H		H		-Me	-CONH-	1

Table 1 (continued)

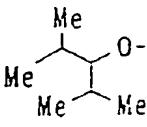
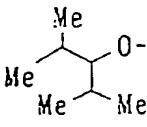
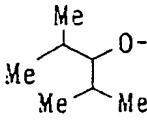
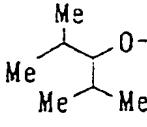
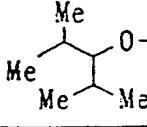
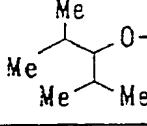
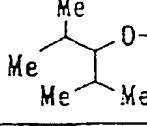
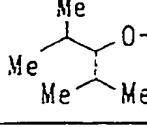
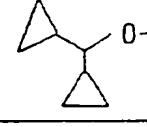
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
521	H		H	H	-Me	-CONMe-	1
522	H		Me-	H	-Me	-CONMe-	1
523	H		Et-	H	-Me	-CONMe-	1
524	H		iPr-	H	-Me	-CONMe-	1
525	H		Me-O-	H	-Me	-CONMe-	1
526	H		Et-O-	H	-Me	-CONMe-	1
527	H		iPr-O-	H	-Me	-CONMe-	1
528	-OH		H	H	-Me	-CONMe-	1
529	H		H	H	-Me	-CONH-	2

Table 1 (continued)

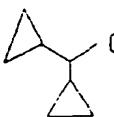
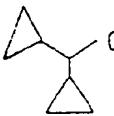
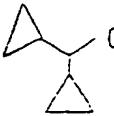
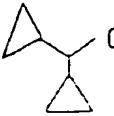
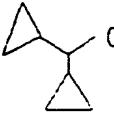
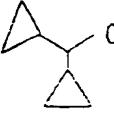
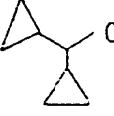
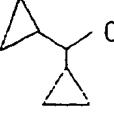
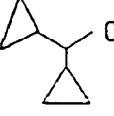
5 Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
10 530	H		Me-	H	-Me	-CONH-	2
15 531	H		Et-	H	-Me	-CONH-	2
20 532	H		iPr-	H	-Me	-CONH-	2
25 533	H		Me-O-	H	-Me	-CONH-	2
30 534	H		Et-O-	H	-Me	-CONH-	2
35 535	H		iPr-O-	H	-Me	-CONH-	2
40 536	-OH		O-	H	H	-CONH-	2
45 537	-Me		O-	H	H	-CONH-	2
50 538	-F		O-	H	H	-CONH-	2

Table 1 (continued)

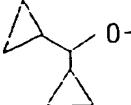
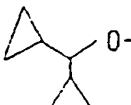
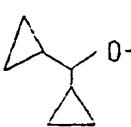
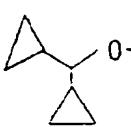
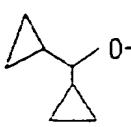
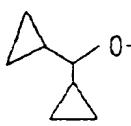
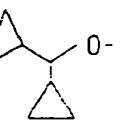
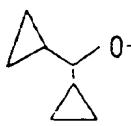
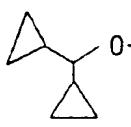
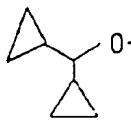
5 Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n	
539	-Cl		O-	H	H	-Me	-CONH-	2
540	-Br		O-	H	H	-Me	-CONH-	2
541	H		O-	H	F-	-Me	-CONH-	2
542	H		O-	H	Cl-	-Me	-CONH-	2
543	H		O-	H	Br-	-Me	-CONH-	2
544	H		O-	H		-Me	-CONH-	2
545	H		O-	H	H	-Me	-CONMe-	2
546	H		O-	Me-	H	-Me	-CONMe-	2
547	H		O-	Et-	H	-Me	-CONMe-	2

Table 1 (continued)

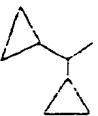
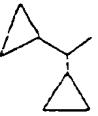
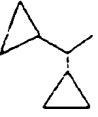
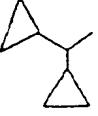
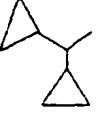
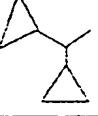
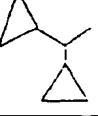
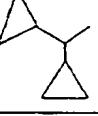
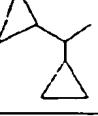
5 Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
10 548	H		O-	iPr-	H	-Me	-CONMe-
15 549	H		O-	Me-O-	H	-Me	-CONMe-
20 550	H		O-	Et-O-	H	-Me	-CONMe-
25 551	H		O-	iPr-O-	H	-Me	-CONMe-
30 552	-OH		O-	H	H	-Me	-CONMe-
35 553	H		O-	H	H	-Me	-CONH-
40 554	H		O-	Me-	H	-Me	-CONH-
45 555	H		O-	Et-	H	-Me	-CONH-
50 556	H		O-	iPr-	H	-Me	-CONH-

Table 1 (continued)

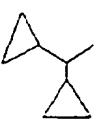
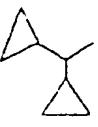
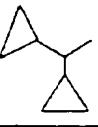
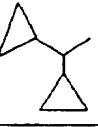
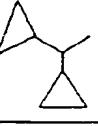
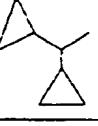
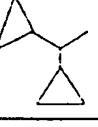
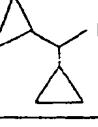
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
557	H		Me-O-	H	-Me	-CONH-	1
558	H		Et-O-	H	-Me	-CONH-	1
559	H		iPr-O-	H	-Me	-CONH-	1
560	-OH		H	H	-Me	-CONH-	1
561	-Me		H	H	-Me	-CONH-	1
562	-F		H	H	-Me	-CONH-	1
563	-Cl		H	H	-Me	-CONH-	1
564	-Br		H	H	-Me	-CONH-	1
565	H		H	F-	-Me	-CONH-	1

Table 1 (continued)

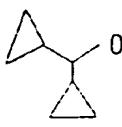
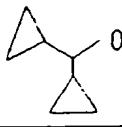
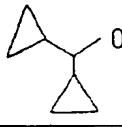
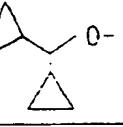
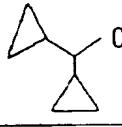
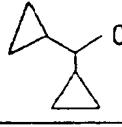
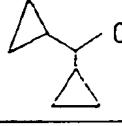
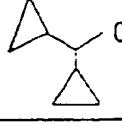
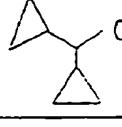
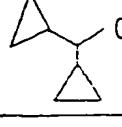
5 Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n	
10 566	H		O-	H	Cl-	-Me	-CONH-	1
15 567	H		O-	H	Br-	-Me	-CONH-	1
20 568	H		O-	H		-Me	-CONH-	1
25 569	H		O-	H	H	-Me	-CONMe-	1
30 570	H		O-	Me-	H	-Me	-CONMe-	1
35 571	H		O-	Et-	H	-Me	-CONMe-	1
40 572	H		O-	iPr-	H	-Me	-CONMe-	1
45 573	H		O-	Me-O-	H	-Me	-CONMe-	1
50 574	H		O-	Et-O-	H	-Me	-CONMe-	1

Table 1 (continued)

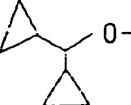
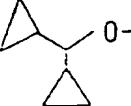
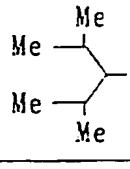
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
575	H		iPr-O-	H	-Me	-CONMe-	1
576	-OH		H	H	-Me	-CONMe-	1
577	H		H	H	-Me	-CONH-	2
578	H		H	H	-Me	-CONH-	2
579	H		H	H	-Me	-CONH-	2
580	H		H	H	-Me	-CONH-	1
581	H		H	H	-Me	-CONH-	2
582	H		H	H	-Me	-CONH-	1
583	H		H	H	-Me	-CONH-	2
584	H		H	H	-Me	-CONH-	2

Table 1 (continued)

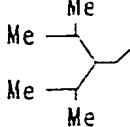
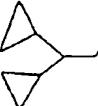
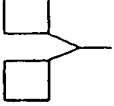
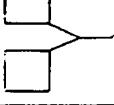
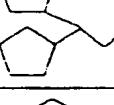
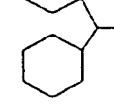
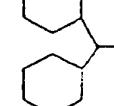
Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
585	H		H	H	-Me	-CONH-	2
586	H		H	H	-Me	-CONH-	2
587	H		H	H	-Me	-CONH-	2
588	H		H	H	-Me	-CONH-	2
589	H		H	H	-Me	-CONH-	2
590	H		H	H	-Me	-CONH-	2
591	H		H	H	-Me	-CONH-	2
592	H		H	H	-Me	-CONH-	2
593	H		H	H	-Me	-CONH-	2

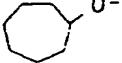
Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
594	H			H	H	-Me	-CONH-
595	H			H	H	-Me	-CONH-
596	H			H	Br-	-Me	-CONH-
597	H		H Me		-Me	-CONH-	2
598	H		-OCH ₂ O-		H	-Me	-CONH-
599	H		-OCH ₂ O-		H	-Me	-CONH-
600	H		-OCH ₂ CH ₂ O-		H	-Me	-CONH-
601	H		-OCH ₂ CH ₂ O-		H	-Me	-CONH-
602	H				H	-Me	-CONH-
603	H				H	-Me	-CONH-
604	H				H	-Me	-CONH-

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n	
605	H			H	-Me	-CONH-	1	
606	H			H	-Me	-CONH-	2	
607	H			H	-Me	-CONH-	1	
608	H			H	-Me	-CONH-	2	
609	H			H	-Me	-CONH-	1	
610	H	t-Bu-		H	t-Bu-	-Me	-CONH-	2
611	H	t-Bu-		H	t-Bu-	-Me	-CONMe-	2
612	H	t-Bu-		H	t-Bu-	-Me	-CONH-	1
613	H	t-Bu-		H	t-Bu-	-Me	-CONMe-	1
614	H	iPr-		H	iPr-	-Me	-CONH-	2
615	H	iPr-		H	iPr-	-Me	-CONH-	1

Table 1 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
616	H	-C(CH ₃) ₂ CH ₂ CH ₂ C(CH ₃) ₂ -		H	-Me	-CONH-	2
617	H	-C(CH ₃) ₂ CH ₂ CH ₂ C(CH ₃) ₂ -		H	-Me	-CONH-	1
618	H	-C(CH ₃) ₂ CH ₂ CH ₂ C(CH ₃) ₂ -		H	-Me	-CONMe-	2
619	H	-C(CH ₃) ₂ CH ₂ CH ₂ C(CH ₃) ₂ -		H	-Me	-CONEt-	2
620	H		H	H	-Me	-SO ₂ NH-	2
621	H		H	H	-Me	-SO ₂ NH-	1
622	H		H	H	-Me	-SO ₂ NMe-	2
623	H		H	H	-Me	-SO ₂ NEt-	2

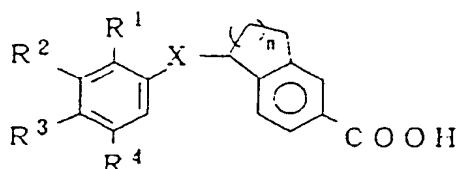
40

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50

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Table 2



Compd. No.	R^1	R^2	R^3	R^4	X	n
624	H		H	H	-CONH-	2
625	H		H	H	-CONH-	1
626	H		H	H	-CONH-	2
627	H		H	H	-CONH-	1
628	H		H	H	-CONH-	2
629	H		H	H	-CONH-	1
630	H		H	H	-CONH-	2
631	H		H	H	-CONH-	1
632	H		H	H	-CONH-	2

Table 2 (continued)

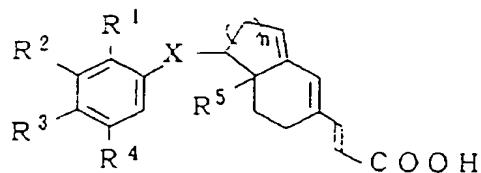
Compd. No.	R ¹	R ²	R ³	R ⁴	X	n
633	H		H	H	-CONH-	1
634	H		H	H	-CONH-	2
635	H		H	H	-CONH-	1
636	H		H	H	-CONH-	2
637	H		H	H	-CONH-	1
638	H		H	H	-CONH-	2
639	H		H	H	-CONH-	2

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Table 3



Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
640	H		H	H	-Me	-CONH-	2
641	H		H	H	-Me	-CONH-	1
642	H		H	H	-Me	-CONH-	2
643	H		H	H	-Me	-CONH-	1
644	H		H	H	-Me	-CONH-	2
645	H		H	H	-Me	-CONH-	1
646	H		H	H	-Me	-CONH-	2
647	H		H	H	-Me	-CONH-	1
648	H		H	H	-Me	-CONH-	2

Table 3 (continued)

Compd. No.	R ¹	R ²	R ³	R ⁴	R ⁵	X	n
649	H		H	H	-Me	-CONH-	1
650	H		H	H	-Me	-CONH-	2
651	H		H	H	-Me	-CONH-	1
652	H		H	H	-Me	-CONH-	2
653	H		H	H	-Me	-CONH-	1
654	H		H	H	-Me	-CONH-	2
655	H		H	H	-Me	-CONH-	2

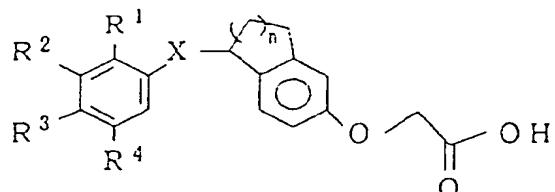
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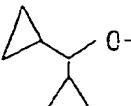
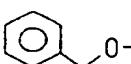
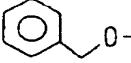
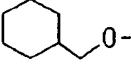
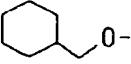
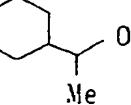
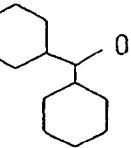
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Table 4



Compd. No.	R^1	R^2	R^3	R^4	X	n
656	H		H	H	-CONH-	2
657	H		H	H	-CONH-	1
658	H		H	H	-CONH-	2
659	H		H	H	-CONH-	1
660	H		H	H	-CONH-	2
661	H		H	H	-CONH-	1
662	H		H	H	-CONH-	2
663	H		H	H	-CONH-	1
664	H		H	H	-CONH-	2

Table 4 (continued)

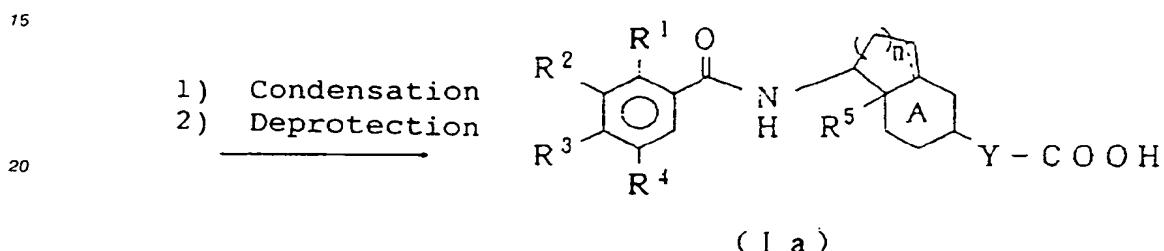
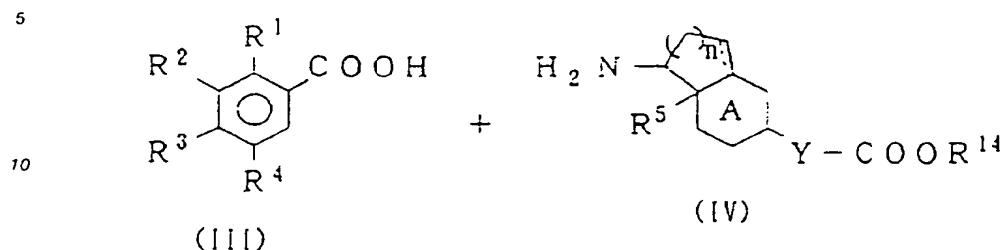
Compd. No.	R ¹	R ²	R ³	R ⁴	X	n
665	H		H	H	-CONH-	1
666	H		H	H	-CONH-	2
667	H		H	H	-CONH-	1
668	H		H	H	-CONH-	2
669	H		H	H	-CONH-	1
670	H		H	H	-CONH-	2
671	H		H	H	-CONH-	2

In Tables 1 - 4, Me means a methyl group, Et means an ethyl group, Pr means a propyl group, iPr means an isopropyl group, Bu means a butyl group, and Ac means an acetyl group.

The compounds of the general formula (I) can form a salt with a base. The base may be selected from those which can form a salt with the compound of the general formula (I). Examples of the salts are metal salts (e.g. sodium salt, magnesium, aluminium salt, etc.), ammonium salt, and amine salts (e.g. methylamine, ethylamine, diethylamine, triethylamine, pyrrolidine, piperidine, morpholine, pyridine, aniline, etc.).

The compounds of the general formula (I) can be prepared, for example, in the manner as shown in the following synthetic route.

(1) Of the compounds of the general formula (I), the compounds wherein X is $\text{-CONR}^{13}\text{-}$ are prepared as follows:

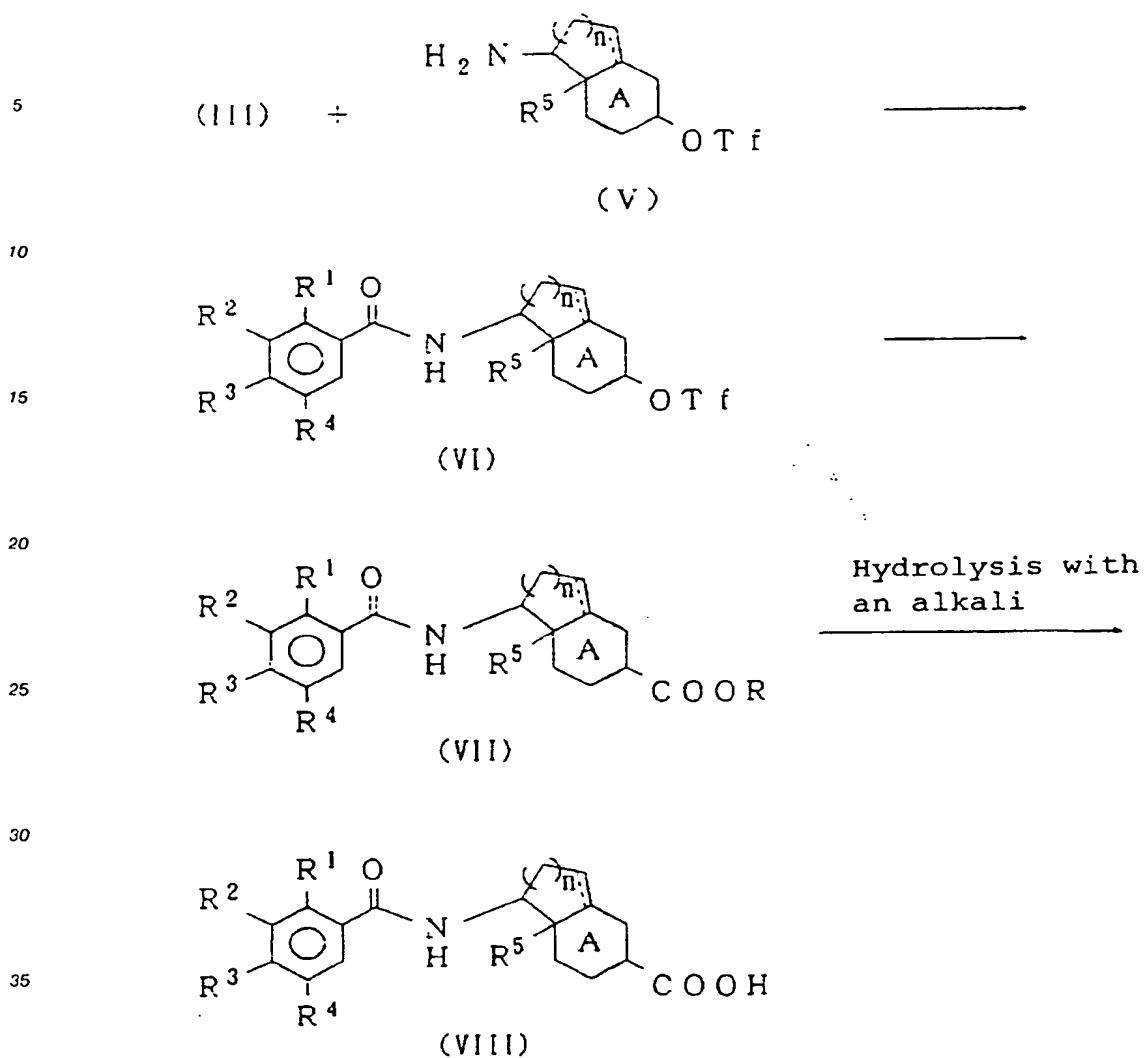


In the above formulae, R¹, R², R³, R⁴, R⁵, n, Y and A have the same significance as defined in the general formula (I) above, and R¹⁴ represents an alkyl group having 1 to 5 carbon atoms.

For example, the benzoic acid derivatives of the general formula (III) is allowed to condense with amino acid derivatives of the general formula (IV) in an appropriate solvent such as methylene chloride, chloroform, tetrahydrofuran, benzene or the like, or without solvent, in the presence or absence of a condensing agent and an organic base, or in the presence of a condensing agent without base. The condensing agent includes inorganic condensing agents such as phosphorus oxychloride, thionyl chloride or the like and organic condensing agents such as dicyclohexylcarbodi-imide, carbodiimidazole, oxalyl chloride, tosyl chloride or the like. The condensed product is subjected to hydrolysis with an alkali to give the objective compound (Ia). The hydrolysis is carried out in a mixture of water and methanol containing an alkali metal hydroxide such as sodium hydroxide or the like, or in a mixture of water, methanol and tetrahydrofuran.

The compound of the general formula (I) in which Y is a single bond can be prepared, for example, by condensing the compound of the general formula (III) with the amine derivative of the general formula (V) below using the method mentioned above, to give the compound of the general formula (VI) below, and this compound is allowed to react with carbon monoxide in the presence of an organic base, phosphine and palladium (II) in an alkanol (ROH) solvent or in a mixture of alkanol (ROH) of 1 -5 carbon atoms with tetrahydrofuran, ether, methylene chloride or the like, to give alkoxycarbonyl compound of the general formula (VII) below, which is then subjected to hydrolysis with an alkali to give the objective compound. (The alkali-hydrolysis is carried out as described above.)

EP 0 579 223 A1



In the above formulae, R¹, R², R³, R⁴, R⁵, n and A have the same significance as defined above, R represents an alkyl group of 1 to 5 alkyl group, and Tf represents trifluoro-methanesulfonyl group.

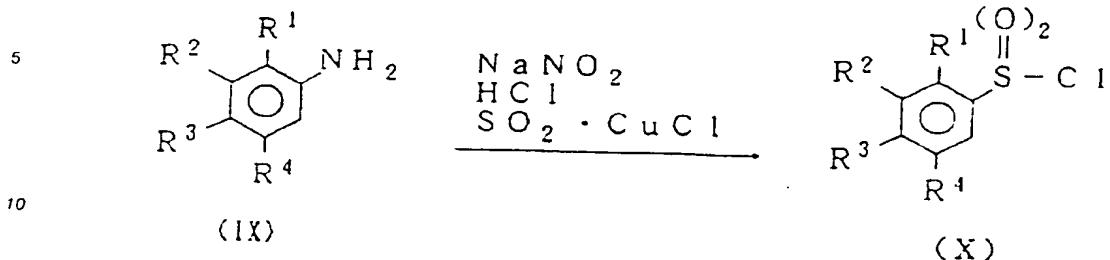
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EP 0 579 223 A1

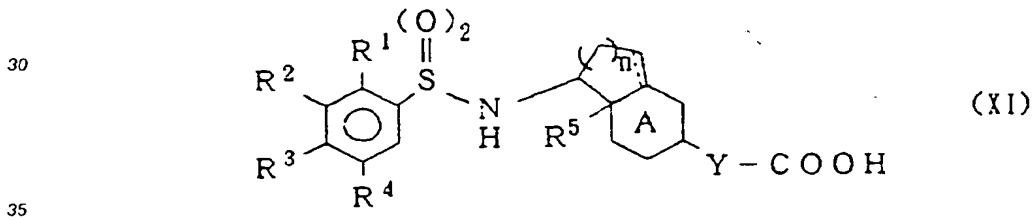
(2) Production of the compound of the general formula (I) in which X is $\text{-SO}_2\text{NR}^{13}$.



In the above formulae, R^1 , R^2 , R^3 and R^4 have the same significance as defined above.

For example, the aniline derivative of the general formula (IX) above is allowed to react with sodium nitrite or potassium nitrite in an appropriate solvent such as aqueous acetic acid, diluted hydrochloric acid or the like, to give a diazonium salt, which is allowed to react with cupric chloride and sulfur dioxide to give the sulfonyl chloride derivative of the general formula (X) above.

The compound of the general formula (XI) below can be prepared by reacting the compound of the general formula (X) with the compound of the general formula (IV) above in the presence or absence of an organic base such as triethylamine, pyridine or the like, in a halogen type solvent such as methylene chloride or the like and subjecting the resultant product to alkali-hydrolysis according to the method as described above.



In the above formulae (XI), R^1 , R^2 , R^3 , R^4 , R^5 , n, Y and A have the same significance as defined in the general formula (I).

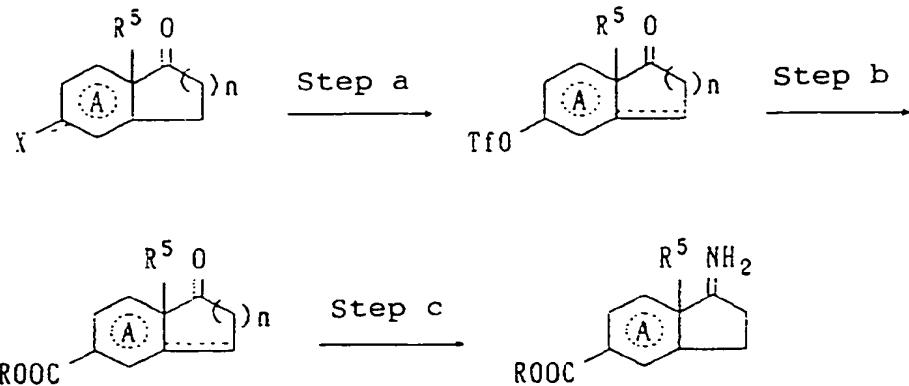
40 (3) Production of amino acid derivatives of the general formula (IV):

i) Production of compounds of the general formula (IV) in which Y is a single bond or $-\text{CH} = \text{CH}-$:

The starting 8a-methyl-3,4,8a-tetrahydro-1,6(2H,7H)-naphthalenedione and 7a-methyl-7,7a-tetrahydro-1,5(6H)-indadione are known compounds, and their optical activity is also known. Furthermore, the starting 6-hydrotetralone and 5-hydroxyindanone are known compounds.

The starting compounds are treated at temperature from -40 °C to room temperature in the presence of 1 mol equivalent of anhydrous trifluoromethanesulfonic acid and tertiary amine such as triethylamine, pyridine or the like, in an inert solvent such as methylene chloride, tetrahydrofuran or the like (Step a), and the resultant product is allowed to react with carbon monoxide or the like in the presence of an organic base, phosphine and palladium (II) in a mixture of alcohol (1 - 5 carbon atoms) with tetrahydrofuran or ether to give an alkoxy compound (Step b). This product is treated with a reducing agent such as sodium cyanoborohydride or the like in the presence of ammonium salt such as ammonium acetate, ammonium chloride or the like, in an alkanol solvent (1 - 5 carbon atoms) or in a mixture of alkanol (1 - 5 carbon atoms) with tetrahydrofuran or ether for subjecting only the ketone group to reductive amination (Step c).

The compound wherein Y is -CH=CH- can be prepared by using an alkyl acrylate or the like in place of carbon monoxide in Step b.



In the above formulae, R⁵, X, n and A have the same significance as defined in the general formula (I), and R represents an alkyl group having 1 - 5 carbon atoms.

20 ii) Production of the compound of the general formula (IV) in which Y is $-OCH_2-$:

For example, 6-hydroxytetralone or 5-hydroxyindanone is allowed to react with α -haloacetic acid ester in the presence of a base such as sodium hydride, triethylamine or the like in a solvent such as methylene chloride, tetrahydrofuran, DMF or the like, and the resultant product

25 is subjected to reductive amination to give the product. The benzoic acid derivative of the formula (III) and aniline derivative of the formula (IX) are commercially available or can be prepared in a conventional manner from commercially available materials.

When the compounds of the present invention are used as medicaments, they can be formulated together with one or more of conventional carriers suitable for desired administering route. For example, formulations such as tablets, capsules, granules, powders, solutions or the like are prepared for oral route. Excipients, binders, lubricants, coloring agents, disintegrators, or the like, which are usually employed for the preparation of pharmaceutical formulations, can be used for preparing solid formulations for oral route. Examples of the excipients are lactose, starch, talc, magnesium stearate, crystalline cellulose, methyl cellulose, carboxymethyl cellulose, glycerin, sodium alginate, arabic gum and the like. Examples of the binders are polyvinyl alcohol, polyvinyl ether, ethyl cellulose, arabic gum, shellac, refined sugar and the like. Examples of the lubricants are-magnesium stearate, talc, and the like. Additionally, coloring agents and disintegrators may be used. Further, tablets may be coated in a conventional manner. Liquid formulations include aqueous or oily suspensions, solutions, syrups, elixirs and the like, and can be prepared in a conventional manner. Injections may be prepared by adding pH adjusting agents, buffers, stabilizers, isotonic agents, topical anesthetics, or the like to the compound of the present invention. Subcutaneous, intramuscular or intravenous injections may be prepared in a conventional manner. For preparing suppositories, oily bases such as cacao oil, polyethylene glycol, Witepsol (Trademark of Dynamite Novel Company) or the like may be used.

Appropriate dosage of the formulation thus obtained varies depending upon symptoms, body weight, age or the like of particular patients. In general, appropriate daily dose to adult is in the range from about 0.01 to 2000 mg, and the daily dose is preferably administered in multiple doses of 1 - 4 times a day.

Appropriate salts of the compound (I) may be prepared using a non-toxic base. Such appropriate salts include those formed with inorganic bases (e.g. sodium salt, potassium salt, etc.), those formed with organic bases (e.g. triethylamine, etc.), and ammonium salts.

EXAMPLES

The present invention will be explained in more detail below by examples and reference examples. The examples are representative only and should not be construed as limiting in any respect.

EP 0 579 223 A1

Example 1

Preparation of 5-(4'-phenoxybenzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 1 in Table 1):

To a solution of 4-phenoxybenzoic acid (410 mg, 1.91 mmol) in methylene chloride (4.0 ml) was added thionyl chloride (0.8 ml), and the resultant mixture was refluxed for 1 hour. The mixture was concentrated, and the residue was dissolved in methylene chloride (4.0 ml) and the resulting solution was dropwise added to a solution of 5-amino-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid methyl ester (420 mg, 1.92 mmol), triethylamine (0.84 ml) and methylene chloride (4.2 ml) under ice cooling. The mixture was stirred at room temperature for 30 minutes, poured into chilled water (50 ml) and extracted with ethyl acetate. The organic extract was concentrated to dryness, and the residue was dissolved in 10% methanolic water (23 ml), mixed with potassium hydroxide (1.09 g, 16.5 mmol) and refluxed for 4 hours. The product was extracted with acidic chloroform, dried over magnesium sulfate and concentrated. The resulting residue was chromatographed on a silica gel column, eluting with chloroform-methanol to give the titled compound (464 mg, yield 60%) as white crystals.

m.p.96-101 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta$ (ppm) = 7.73(d, 2H, 8.7Hz), 7.36(t-like, 2H, 7.4Hz), 7.1-7.2(m, 2H), 6.9-7.1(m, 4H), 5.96(br-d, 1H, 8.7Hz), 5.90(br-s, 1H), 4.10-4.25(m, 1H), 4.10-4.25(m, 1H), 2.10-2.60(m, 4H), 1.75-1.95(m, 3H), 1.40-1.55(m, 1H), 1.01(s, 1H)

IR(KBr): ν (cm $^{-1}$) = 3296, 2938, 1674, 1634, 1588, 1543, 1489, 1244, 1169.

Example 2

Preparation of 5-(3'-phenoxybenzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 10 in Table 1):

The reaction was effected with 3-phenoxybenzoic acid (542 mg, 2.53 mmol) in the same manner as in Example 1 to give the titled compound (412 mg, yield 40%).

m.p.181-186 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta$ (ppm) = 7.33-7.49(m, 5H), 7.10-7.20 (m, 5H), 5.98(d, 1H, J = 9.7Hz), 5.91(t, 1H, J = 3.7Hz), 4.13-4.26(m, 1H), 2.14-2.59(m, 3H), 1.82-1.92(m, 4H), 1.40-1.53(m, 1H), 1.02(s, 3H)

IR(KBr): ν (cm $^{-1}$) = 3285, 3069, 9240, 1672, 1634, 1580, 1555, 1483, 1427, 1306, 1275, 1233.

Example 3

Preparation of 5-(2'-phenoxybenzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 12 in Table 1):

The reaction was effected with 2-phenoxybenzoic acid (488 mg, 2.28 mmol) as in the same manner as in Example 1 to give the titled compound (330 mg, yield 36%).

m.p.75-77 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta$ = 8.23(dd, 1H, 1.8Hz, 7.9Hz), 7.73(br-d, 1H, 8.0Hz), 7.3-7.42(m, 3H), 7.1-7.3(m, 3H), 7.00(d, 2H, 7.9Hz), 6.87(d, 1H, 8.2Hz), 5.85(br-s, 1H), 4.1-4.2(m, 1H), 2.2-2.5(m, 3H), 2.0-2.2(m, 1H), 1.8-1.9(m, 1H), 1.6-1.8(m, 2H), 1.2-1.4(m, 1H), 0.84(s, 3H)

IR(KBr): ν (cm $^{-1}$) = 3403, 3069, 2932, 1640, 1601, 1534, 1478, 1449, 1223, 753.

Example 4

Preparation of 5-[3'-(p-isopropylphenoxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 16 in Table 1):

The reaction was effected with 3-(p-isopropylphenoxy)benzoic acid (177 mg, 0.74 mmol) in the same manner as in Example 1 to give the titled compound (253 mg, yield 77%). m.p.146-149 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta$ = 7.3-7.45(m, 3H), 7.15-7.23(m, 3H), 7.10(dd, 1.4Hz, 8.0Hz), 6.90-6.96(m, 2H), 5.98(br-d, 1H, 8.7Hz), 5.90(br-s, 1H), 4.1-4.2(m, 1H), 2.80-2.95(m, 1H), 2.10-2.60(m, 4H), 1.70-1.95(m, 3H), 1.35-1.50(m, 1H), 1.22, 1.25(each s, each 3H), 0.99(s, 3H)

IR(KBr): ν (cm $^{-1}$) = 3312, 2959, 2930, 2870, 1674, 1636, 1582, 1541, 1507, 1480, 1429, 1306, 1279, 1236.

Example 5

Preparation of 5-[3'-(m-isopropylphenoxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 17 in Table 1):

5 The reaction was effected with 3-(m-isopropylphenoxy)benzoic acid (704 mg, 2.75 mmol) in the same manner as in Example 1 to give the titled compound (612 mg, yield 50%).

m.p.83-86 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.3\text{-}7.47(\text{m}, 3\text{H}), 7.25(\text{dd}, 1\text{H}, 7.9\text{Hz}, 7.8\text{Hz}), 7.17(\text{br-s}, 1\text{H}), 7.10(\text{ddd},$

10 1.3Hz, 2.2Hz, 7.9Hz), 7.00(br-d, 1H, 7.8Hz), 6.90(dd, 1H, 2.1Hz, 1.8Hz), 6.79(ddd, 1H, 0.8Hz, 2.2Hz, 7.9Hz), 5.95(d, 1H, 9.7Hz), 5.90(dd, 1H, 3.8Hz, 3.7Hz), 4.11-4.23 (m, 1H), 2.87(dq, 1H, each 6.9Hz), 2.52(dd, 1H, 4.7Hz, 19Hz), 2.10- 2.47(m, 3H), 1.76-1.90(m, 3H), 1.37-1.49(m, 1H), 1.22(d, 6H, 6.9Hz), 1.02(s, 3H)

IR(KBr): $\nu(\text{cm}^{-1}) = 3285, 3071, 2961, 2870, 2822, 1672, 1634, 1576, 1555, 1481, 1427, 1308, 1273, 1244.$

Example 6

Preparation of 5-[3'-(o-isopropylphenoxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 18 in Table 1):

20 The reaction was effected with 3-(o-isopropylphenoxy)benzoic acid (347 mg, 1.44 mmol) in the same manner as in Example 1 to give the titled compound (344 mg, yield 54%).

m.p.192-194 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.3\text{-}7.4(\text{m}, 4\text{H}), 7.1\text{-}7.2(\text{m}, 2\text{H}), 6.98(\text{dd-like}, 1\text{H}, 1.4\text{Hz}, 8.0\text{Hz}), 6.80\text{-}6.90(\text{m},$

25 1H), 5.98(br-d, 1H, 8.7Hz), 5.90(br-s, 1H), 4.1-4.25(m, 1H), 3.1-3.3(m, 1H), 2.1-2.6 (m, 4H), 1.7-1.9(m, 3H)

IR(KBr): $\nu(\text{cm}^{-1}) = 3291, 3071, 2961, 2870, 2824, 2629, 1674, 1634, 1578, 1555, 1483, 1451, 1429, 1306, 1275, 1235, 1184.$

Example 7

30 Preparation of 5-[3'-(p-tolyloxy)benzoylamino]-10-methyl- $\Delta^{1(2), 8(9)}$ -octalin-2-carboxylic acid (Compound No. 21 in Table 1):

To a mixture of 3-(p-tolyloxy)benzoic acid (414 mg, 1.81 mmol), 5-amino-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-35 carboxylic acid methyl ester (401 mg, 1.81 mmol), triethylamine (1.26 mg, 9.05 mmol) and methylene chloride (4.1 ml) was added thionyl chloride (396 μl , 5.43 mmol) with ice cooling. The resultant mixture was stirred at room temperature for 30 minutes, poured into chilled water (50 ml), and extracted with ethyl acetate. The organic extract was concentrated, and the residue was dissolved in a mixture of 10% methanol (12 ml) and THF (3.7 ml), mixed with potassium hydroxide (566 mg, 8.57 mmol) and refluxed for 3 hours.

40 The product was extracted with acidic chloroform, and the organic layer was dried and concentrated. The resultant residue was chromatographed on a silica gel column, eluting with chloroform-methanol to give the titled compound (263 mg, yield 35%).

m.p.168-172 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.3\text{-}7.4(\text{m}, 3\text{H}), 7.0\text{-}7.2(\text{m}, 4\text{H}), 6.90 (\text{d}, 2\text{H}, 8.5\text{Hz}), 6.00(\text{d}, 1\text{H}, 9.6\text{Hz}), 5.89\text{-}$

45 (br-s, 1H), 4.1-4.2(m, 1H), 2.1-2.6(m, 4H), 2.32(s, 3H), 1.7-1.9(m, 3H), 1.40(ddd, 1H, 5.7Hz, 5.7Hz, 13Hz), 0.99(s, 3H)

IR(KBr): $\nu(\text{cm}^{-1}) = 3283, 2940, 1672, 1634, 1580, 1559, 1507, 1481, 1427, 1306, 1275, 1238, 1209, 1186.$

Example 8

50 Preparation of 5-[3'-(p-t-butylphenoxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylicacid (Compound No. 24 in Table 1):

The reaction was effected with 3-(p-t-butylphenoxy)benzoic acid (435 mg, 1.61 mmol) in the same manner as in Example 7 to give the titled compound (282 mg, yield 38%).

m.p.233-238 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.3\text{-}7.45(\text{m}, 5\text{H}), 7.16(\text{d}, 1\text{H}, 1.5\text{Hz}), 7.52(\text{dd-like}, 1\text{H}, 1\text{Hz}, 1.5\text{Hz}, 8.0\text{Hz}), 6.94(\text{d}, 2\text{H}, 8.9\text{Hz}), 5.96 (\text{br-d}, 1\text{H}, 8.7\text{Hz}), 5.86(\text{br-s}, 1\text{H}), 4.05\text{-}4.20(\text{m}, 1\text{H}), 2.10\text{-}2.60(\text{m}, 4\text{H}), 1.75\text{-}1.95(\text{m},$

3H), 1.40-1.55(m, 1H), 1.30(s, 9H), 0.99(s, 3H)

IR(KBr): ν (cm⁻¹) = 3266, 2965, 2870, 1680, 1634, 1613, 1580, 1543, 1508, 1480, 1426, 1306, 1279, 1236.

Example 9

Preparation of 5-{3''-[p-(2'-methylpropyl)phenoxy]benzoylamino}-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 30 in Table 1):

The reaction was effected with 3-p-(2'-methylpropyl)phenoxybenzoic acid (246 mg, 0.91 mmol) in the same manner as in Example 7 to give the titled compound (290 mg, yield 69%).

m.p.142-148 °C

¹H-NMR(CDCl₃, 250MHz) δ = 7.3-7.45(m, 3H), 7.18(s-like, 1H), 7.06-7.13(m, 3H), 6.92(d, 2H, 11.3Hz), 5.95(d, 1H, 9.7Hz), 5.90 (br-s, 1H), 4.10-4.22(m, 1H), 4.52(dd, 1H, 4.1Hz, 18.3Hz), 2.44(d, 2H, 7.2Hz), 2.15-4.60(m, 3H), 1.75-1.95(m, 4H), 1.42(ddd, 1H, 5.7Hz, 5.7Hz, and 13Hz), 0.99(s, 3H), 0.89(d, 6H, 6.5Hz)

IR(KBr): ν (cm⁻¹) = 3289, 2959, 2870, 1672, 1638, 1609, 1584, 1545, 1505, 1481, 1431, 1308, 1279, 1235.

Example 10

Preparation of 5-[3'-(p-cyclopentylphenoxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 33 in Table 1):

The reaction was effected with 3-(p-cyclopentylphenoxy)benzoic acid (488 mg, 1.73 mmol) in the same manner as in Example 7 to give the titled compound (240 mg, yield 30%).

m.p.229-231 °C

¹H-NMR(CDCl₃, 250MHz) δ = 7.3-7.4(m, 3H), 7.15-7.25(m, 3H), 7.09(ddd, 1H, 1.4Hz, 0, 8Hz, 9.2Hz), 6.93(d, 2H, 8.5Hz), 5.96(d, 1H, 9.6Hz), 5.89(br-s, 1H), 4.05-4.20(m, 1H), 2.85-3.05(m, 1H), 1.95-2.30(m, 6H), 1.30-1.90(m, 10H), 0.99(s, 3H)

IR(KBr): ν (cm⁻¹) = 3337, 2944, 2868, 1676, 1638, 1613, 1578, 1543, 1505, 1483, 1426, 1323, 1304, 1275, 1236.

Example 11

Preparation of 5-[3'-(p-cyclohexylphenoxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 36 in Table 1):

The reaction was effected with 3-(p-cyclohexylphenoxy)benzoic acid (415 mg, 1.40 mmol) in the same manner as in Example 7 to give the titled compound (263 mg, yield 33%).

m.p.124-125 °C

¹H-NMR(CDCl₃, 250MHz) δ = 7.3-7.45(m, 3H), 7.1-7.2(m, 3H), 7.10(ddd, 1H, 1.3Hz, 2.4Hz, 7.9Hz), 6.92(d, 2H, 8.5Hz), 5.97(br-d, 1H, 9.5Hz), 5.89(br-s, 1H), 4.10-4.25(m, 1H), 2.10-2.60(m, 5H), 1.65-1.95(m, 8H), 1.10-1.50(m, 6H), 0.99(s, 3H)

IR(KBr): ν (cm⁻¹) = 3443, 2926, 2851, 1676, 1636, 1580, 1541, 1507, 1480, 1429, 1306, 1273, 1238.

Example 12

Preparation of 5-[3'-(p-N-methylaminocarbonylphenoxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 38 in Table 1):

The reaction was effected with 3-(p-N-methyl-aminocarbonylphenoxy)benzoic acid (448 mg, 1.19 mmol) in the same manner as in Example 7 to give the titled compound (35 mg, yield 6.4%).

m.p.123-127 °C

¹H-NMR(CDCl₃, 250MHz) δ = 7.72(d, 2H, 8.6Hz), 7.35-7.55(m, 3H), 7.10-7.20(m, 2H), 6.97(d, 2H, 8.6Hz), 6.30(br-d-like), 6.10(br-d, 1H, 9.5Hz), 5.87(br-s, 1H), 2.96, 2.98(each s, total 3H), 2.1-2.6 (m, 4H), 1.6-1.9(m, 3H), 1.3-1.5(m, 1H), 0.99(s, 3H)

IR(KBr): ν (cm⁻¹) = 3337, 3069, 2936, 1684, 1636, 1578, 1543, 1501, 1316, 1240, 1177.

EP 0 579 223 A1

Example 13

Preparation of 5-[3'-(2', 4-dimethylphenoxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 42 in Table 1):

The reaction was effected with 3-(2',4'-dimethylphenoxy) benzoic acid (271 mg, 1.12 mmol) in the same manner as in Example 7 to give the titled compound (135 mg, yield 28%).

m.p.137-142 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.3\text{-}7.45(\text{m}, 3\text{H}), 7.18(\text{d}, 1\text{H}, 2.0\text{Hz}), 7.09(\text{ddd}, 1\text{H}, 1.2\text{Hz}, 2.2\text{Hz}, 9.0\text{Hz}),$

6.76(d, 1H, 0.6Hz), 6.62(d, 2H, 0.6Hz), 5.97(br-d, 1H, 9.6Hz), 5.89(dd, 1H, each 3.7Hz), 4.10-4.25 (m, 1H), 2.1-2.6(m, 4H), 2.27(s, 6H), 2.7-2.95(m, 3H), 1.43(ddd, 1H, 5.7Hz, 5.7Hz and 13Hz), 0.99(s, 3H)

IR(KBr): $\nu(\text{cm}^{-1}) = 3290, 2938, 1676, 1636, 1578, 1541, 1476, 1456, 1296, 1275, 1225, 1142.$

Example 14

Preparation of 5-[3'-(5'',6'',7'',8''-tetrahydro-2''-naphthoxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 45 in Table 1):

The reaction was effected with 3-(5',6',7',8'-tetrahydro-2'-naphthoxy)benzoic acid (324 mg, 1.21 mmol) in the same manner as in Example 7 to give the titled compound (131 mg, yield 24%).

m.p.141-151 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.30\text{-}7.50(\text{m}, 3\text{H}), 7.20(\text{br-s}, 1\text{H}), 7.11(\text{ddd}, 1\text{H}, 1.5\text{Hz}, 2.5\text{Hz}, 7.8\text{Hz}), 7.04\text{-}(d, 1\text{H}, 8.1\text{Hz}), 6.70\text{-}6.80 (\text{m}, 2\text{H}), 5.97(d, 1\text{H}, 9.7\text{Hz}), 5.92(\text{br-s}, 1\text{H}), 4.12\text{-}4.25(\text{m}, 1\text{H}), 2.7\text{-}2.8(\text{m}, 4\text{H}), 2.54\text{-}(\text{dd}, 1\text{H}, 4.7\text{Hz}, 18.1\text{Hz}), 2.2\text{-}2.5(\text{m}, 3\text{H}), 1.70\text{-}2.0 (\text{m}, 7\text{H}), 1.45(\text{ddd}, 1\text{H}, 5.2\text{Hz}, 5.2\text{Hz and } 12.5\text{Hz}), 1.02(\text{s}, 3\text{H})$

IR(KBr): $\nu(\text{cm}^{-1}) = 3287, 2934, 2859, 2633, 1674, 1636, 1580, 1557, 1497, 1481, 1427, 1306, 1275, 1248, 1188, 1146.$

Example 15

Preparation of 5-(3'-benzyloxy-benzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 46 in Table 1):

The reaction was effected with 3-benzyloxybenzoic acid (516 mg, 2.26 mmol) in the same manner as in Example 1 to give the titled compound (325 mg, yield 34.5%).

m.p.175-177 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.30\text{-}7.47(\text{m}, 8\text{H}), 7.21(\text{s}, 1\text{H}), 7.13 (\text{dd}, 1\text{H}, 2.5\text{Hz}, 8.0\text{Hz}), 5.99(\text{d}, 1\text{H}, J = 9.7\text{Hz}), 5.92(\text{t}, 1\text{H}, J = 3.7\text{Hz}), 5.13(\text{s}, 2\text{H}), 4.16\text{-}4.26(\text{m}, 1\text{H}), 2.31\text{-}2.60(\text{m}, 4\text{H}), 1.80\text{-}1.92(\text{m}, 3\text{H}), 1.47(\text{dt}, 1\text{H}, J = 5.5\text{Hz}, 12.8\text{Hz})$

IR(KBr): $\nu(\text{cm}^{-1}) = 3291, 3065, 3036, 2942, 2910, 2870, 1680, 1640, 1611, 1580, 1549, 1483, 1454, 1426,$

1304, 1285, 1235.

Example 16

Preparation of 5-(3'-diphenylmethoxybenzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 50 in Table 1):

The reaction was effected with 3-diphenylmethoxybenzoic acid (498 mg, 1.64 mmol) in the same manner as in Example 1 to give the titled compound (204 mg, yield 25%).

m.p.89-91 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.1\text{-}7.5(\text{m}, 14\text{H}), 7.0\text{-}7.1(\text{m}, 1\text{H}), 6.25(\text{s}, 1\text{H}), 5.8\text{-}5.9(\text{m}, 2\text{H}), 4.0\text{-}4.2(\text{m}, 1\text{H}), 2.1\text{-}2.6(\text{m}, 4\text{H}), 2.7\text{-}2.9(\text{m}, 3\text{H}), 1.35\text{-}1.5(\text{m}, 1\text{H}), 0.96(\text{s}, 3\text{H})$

IR(KBr): $\nu(\text{cm}^{-1}) = 3308, 3063, 3030, 2934, 1684, 1634, 1580, 1524, 1483, 1454, 1429, 1273, 1233, 1186, 1020, 748, 700.$

EP 0 579 223 A1

Example 17

Preparation of 5-(3'-cyclopentyloxybenzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 51 in Table 1):

5 The reaction was effected with 3-cyclopentyloxybenzoic acid (413 mg, 2.0 mmol) in the same manner as in Example 1 to give the titled compound (230 mg, yield 29%).

m.p.175-176 °C

10 $^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.26\text{-}7.36(\text{m}, 3\text{H}), 7.21(\text{s}, 1\text{H}), 7.02(\text{dd}, 1\text{H}, J = 2.5\text{Hz}, 8.0\text{Hz}), 6.01(\text{d}, 1\text{H}, J = 9.7\text{Hz}), 5.92(\text{t}, 1\text{H}, J = 3.7\text{Hz}), 4.80\text{-}4.88(\text{m}, 1\text{H}), 4.16\text{-}4.26(\text{m}, 1\text{H}), 2.22\text{-}2.59(\text{m}, 3\text{H}), 1.56\text{-}2.00(\text{m}, 12\text{H}), 1.46(\text{dt}, 1\text{H}, J = 5.5\text{Hz}, 12.8\text{Hz}), 1.03(\text{s}, 3\text{H})$

IR(KBr): $\nu(\text{cm}^{-1}) = 3291, 2936, 2870, 1680, 1636, 1589, 1555, 1485, 1426, 1319, 1275, 1242.$

Example 18

15 Preparation of 5-(3'-cyclohexyloxybenzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 63 in Table 1):

20 The reaction was effected with 3-cyclohexyloxybenzoic acid (530 mg, 2.41 mmol) in the same manner as in Example 1 to give the titled compound (292 mg, yield 29.0%).

m.p.192-194 °C

25 $^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.21\text{-}7.37(\text{m}, 4\text{H}), 7.04(\text{dd}, 1\text{H}, 2.5\text{Hz}, 8.0\text{Hz}), 6.01(\text{d}, 1\text{H}, J = 9.3\text{Hz}), 5.92(\text{t}, 1\text{H}, J = 3.7\text{Hz}), 4.15\text{-}4.38 (\text{m}, 2\text{H}), 2.21\text{-}2.59(\text{m}, 4\text{H}), 1.76\text{-}2.05(\text{m}, 7\text{H}), 1.29\text{-}1.64(\text{m}, 7\text{H}), 1.03(\text{s}, 3\text{H})$

IR(KBr): $\nu(\text{cm}^{-1}) = 3312, 2936, 2865, 1672, 1636, 1609, 1576, 1539, 1481, 1429, 1304, 1287, 1236, 1051.$

Example 19

Preparation of 5-(3'-cycloheptyloxybenzoylamino)-10-methy- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 111 in Table 1):

30 The reaction was effected with 3-cycloheptyloxybenzoic acid (2.17 mg, 9.26 mmol) in the same manner as in Example 7 to give the titled compound (2.8 mg, yield 71%).

m.p.224-226 °C

35 $^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.1\text{-}7.4(\text{m}, 4\text{H}), 6.98\text{-}7.02(\text{m}, 1\text{H}), 6.01(\text{br-d}, 1\text{H}, 9.7\text{Hz}), 5.92(\text{t-like}, 1\text{H}, J = 1.0\text{Hz}), 4.20\text{-}4.35(\text{m}, 1\text{H}), 4.1\text{-}4.3(\text{m}, 1\text{H}), 2.1\text{-}2.6(\text{m}, 4\text{H}), 2.0\text{-}2.1(\text{m}, 2\text{H}), 1.3\text{-}1.9(\text{m}, 8\text{H}), 1.03(\text{s}, 3\text{H})$

IR(KBr): $\nu(\text{cm}^{-1}) = 3304, 2932, 2863, 1674, 1638, 1609, 1576, 1545, 1483, 1456, 1427, 1323, 1306, 1277, 1236, 1188, 1022, 750.$

Example 20

40 Preparation of 5-(3'-cyclooctyloxybenzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 159 in Table 1):

45 The reaction was effected with 3-cyclooctyloxybenzoic acid (268 mg, 1.08 mmol) in the same manner as in Example 7 to give the titled compound (358 mg, yield 76%).

m.p.214-218 °C

50 $^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.2\text{-}7.35(\text{m}, 3\text{H}), 7.17(\text{d}, 1\text{H}, 1.4\text{Hz}), 7.00(\text{dd}, 1\text{H}, 2.4\text{Hz}, 8.0\text{Hz}), 5.98(\text{d}, 1\text{H}, 10\text{Hz}), 5.90(\text{t}, 1\text{H}, 3.6\text{Hz}), 4.4\text{-}4.5(\text{m}, 1\text{H}), 4.18(\text{ddd}, 1\text{H}, 5.2\text{Hz}, 10\text{Hz and } 19\text{Hz}), 2.53(\text{dd}, 1\text{H}, 5.6\text{Hz}, 19\text{Hz}), 2.1\text{-}2.4(\text{m}, 3\text{H}), 1.4\text{-}2.0(\text{m}, 18\text{H}), 1.01(\text{s}, 3\text{H})$

55 IR(KBr): $\nu(\text{cm}^{-1}) = 3299, 2928, 1676, 1638, 1609, 1576, 1549, 1478, 1424, 1323, 1287.$

EP 0 579 223 A1

Example 21

Preparation of 5-[3'-(4-methylcyclohexyloxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 207 in Table 1):

5 The reaction was effected with 3-(4-methylcyclohexyloxy) benzoic acid (423 mg, 1.81 mmol) in the same manner as in Example 1 to give the titled compound (375 mg, yield 49.0%).

m.p.194-199 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.21\text{-}7.40(\text{m}, 4\text{H}), 7.06(\text{dd}, 1\text{H}, J=2.5\text{Hz}, 8.0\text{Hz}), 5.92(\text{t}, 1\text{H}, J=3.7\text{Hz}),$

10 5.51(d, 1H, J=9.7Hz), 4.14-4.61(m, 2H), 2.23-2.60(m, 3H), 1.34-2.07(m, 14H), 1.03 (s, 3H), 0.94(d, 3H, J=5.0Hz)

IR(KBr): $\nu(\text{cm}^{-1}) = 3293, 2932, 2870, 2633, 1674, 1637, 1578, 1545, 1481, 1427, 1277, 1236, 1124, 1036.$

Example 22

15 Preparation of 5-[3-(4-isopropylcyclohexyloxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 255 in Table 1):

The reaction was effected with 3-(4-isopropylcyclohexyloxy)benzoic acid (474 mg, 1.81 mmol) in the same manner as in Example 1 to give the titled compound (185 mg, yield 23%).

m.p.128-129 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.20\text{-}7.40(\text{m}, 4\text{H}), 7.10(\text{s-like}, 1\text{H}), 6.95\text{-}7.05(\text{m}, 1\text{H}), 5.99(\text{br-d}, 1\text{H}, 9.8\text{Hz}), 5.86(\text{t-like}, 1\text{H}, J=1\text{Hz}), 4.1\text{-}4.3(\text{m}, 2\text{H}), 2.1\text{-}2.6(\text{m}, 6\text{H}), 1.8\text{-}2.0(\text{m}, 5\text{H}), 1.1\text{-}1.6(\text{m}, 7\text{H}), 1.02(\text{s}, 3\text{H}), 0.88(\text{s}, 6\text{H})$

25 IR(KBr): $\nu(\text{cm}^{-1}) = 3324, 2938, 2868, 2627, 1678, 1636, 1580, 1535, 1483, 1431, 1275, 1238, 1132, 1080, 1040, 1003.$

Example 23

30 Preparation of 5-[3-(4-trans-t-butylcyclohexyloxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 303 in Table 1):

The reaction was effected with 3-(4-trans-t-butylcyclohexyloxy)benzoic acid (237 mg, 0.86 mmol) in the same manner as in Example 1 to give the titled compound (154 mg, yield 38.5%).

35 m.p.129-134 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.21\text{-}7.39(\text{m}, 4\text{H}), 7.04(\text{ddd}, 1\text{H}, J=2.5\text{Hz}, 2.5\text{Hz}, 8.0\text{Hz}), 6.03(\text{d}, 1\text{H}, 9.7\text{Hz}), 5.92(\text{t}, 1\text{H}, J=3.7\text{Hz}), 4.14\text{-}4.29(\text{m}, 1\text{H}), 2.15\text{-}2.61(\text{m}, 6\text{H}), 1.82\text{-}1.96(\text{m}, 5\text{H}), 1.31\text{-}1.54 (\text{m}, 3\text{H}), 1.09\text{-}1.18(\text{m}, 3\text{H}), 1.03(\text{s}, 3\text{H}), 0.88(\text{s}, 9\text{H})$

IR(KBr): $\nu(\text{cm}^{-1}) = 3439, 2946, 2866, 1678, 1638, 1582, 1528, 1481, 1275, 1236, 1049, 1030.$

40 **Example 24**

Preparation of 5-[3'-(4"-cis-t-butylcyclohexyloxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 351 in Table 1):

45 The reaction was effected with 3-(4-cis-t-butylcyclohexyloxy)benzoic acid (320 mg, 1.16 mmol) in the same manner as in Example 1 to give the titled compound (102 mg, yield 52.6%).

m.p.137-139 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.21\text{-}7.37(\text{m}, 4\text{H}), 7.06(\text{ddd}, 1\text{H}, J=2.5\text{Hz}, 2.5\text{Hz}, 8\text{Hz}), 5.92(\text{t}, 1\text{H},$

50 J=3.7Hz), 4.61(br s, 1H), 4.16-4.27 (m, 1H), 2.06-2.62(m, 7Hz), 1.81-1.96(m, 3H), 1.39-1.62(m, 7H), 1.03(s, 3H), 0.88(s, 9H)

IR(KBr): $\nu(\text{cm}^{-1}) = 3319, 2943, 2868, 1676, 1636, 1582, 1530, 1481, 1429, 1306, 1273, 1236, 1182, 1007.$

EP 0 579 223 A1

Example 25

Preparation of 5-[3'-(N-acetyl-3'-piperidinyloxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 299 in Table 1):

5 The reaction was effected with 3-(N-acetyl-3'-piperidinyloxy)benzoic acid (203 mg, 0.77 mmol) in the same manner as in Example 7 to give the titled compound (303 mg, yield 87%).
m.p.68-72 °C

10 $^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.40(\text{d-like}, 1\text{H}), 7.35(\text{d}, 1\text{H}, 7.9\text{Hz}), 7.25\text{-}7.29(\text{m}, 1\text{H}), 7.19(\text{d}, 1\text{H}, 1.5\text{Hz}), 7.06(\text{ddd}, 1\text{H}, 1.6\text{Hz}, 0.8\text{Hz}, 7.1\text{Hz}), 6.02(\text{br-d}, 1\text{H}, 9.7\text{Hz}), 5.93(\text{t}, 1\text{H}, 3.7\text{Hz}), 4.61\text{-}4.67 (\text{m}, 1\text{H}), 4.14\text{-}4.23(\text{m}, 1\text{H}), 3.3\text{-}3.9(\text{m}, 4\text{H}), 2.1\text{-}2.6(\text{m}, 4\text{H}), 2.13 (\text{s}, 1\text{H}), 1.7\text{-}2.0(\text{m}, 3\text{H}), 1.4\text{-}1.6(\text{m}, 1\text{H}), 1.04(\text{s}, 3\text{H})$
IR(KBr): $\nu(\text{cm}^{-1}) = 3384, 2934, 1698, 1632, 1580, 1539, 1483, 1454, 1364, 1318, 1233, 1034$.

Example 26

15 Preparation of 5-[3'-(N-methyl-3'-piperidinyloxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 401 in Table 1):

20 The reaction was effected with 3-(N-methyl-3'-piperidinyloxy)benzoic acid (168 mg, 0.62 mmol) in the same manner as in Example 7 to give the titled compound (70 mg, yield 26%).

m.p.63-65 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.35\text{-}7.45(\text{m}, 3\text{H}), 7.11(\text{d}, 1\text{H}, 1.9\text{Hz}), 7.08(\text{ddd}, 1\text{H}, 2.7\text{Hz}, 2.5\text{Hz}, 6.5\text{Hz}), 6.84(\text{br-d}, 1\text{H}, 9.6\text{Hz}), 5.88 (\text{br-s}, 1\text{H}), 4.75(\text{br-s}, 1\text{H}), 4.1\text{-}4.3(\text{m}, 1\text{H}), 3.2\text{-}3.4(\text{m}, 4\text{H}), 2.82 (\text{s}, 3\text{H}), 1.8\text{-}2.6(\text{m}, 11\text{H}), 1.4\text{-}1.6(\text{m}, 1\text{H}), 1.04(\text{s}, 3\text{H})$

25 IR(KBr): $\nu(\text{cm}^{-1}) = 3420, 2938, 2722, 1692, 1640, 1582, 1537, 1481, 1235, 1040, 754, 691$.

Example 27

30 Preparation of 5-[3'-(3''-tetrahydropyranyl)oxybenzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 403 in Table 1):

The reaction was effected with 3-(3'-tetrahydropyranyl) oxybenzoic acid (313 mg, 1.41 mmol) in the same manner as in Example 7 to give the titled compound (229 mg, yield 39%).

m.p.83-85 °C

35 $^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.39(\text{d}, 1\text{H}, 1.4\text{Hz}), 7.34(\text{d}, 1\text{H}, 7.8\text{Hz}), 7.20(\text{s-like}, 1\text{H}), 7.06(\text{ddd}, 1\text{H}, 1.3\text{Hz}, 1.1\text{Hz}, 8.2\text{Hz}), 6.03(\text{d}, 1\text{H}, 9.6\text{Hz}), 5.92(\text{s-like}, 1\text{H}), 4.5\text{-}4.7(\text{m}, 1\text{H}), 4.1\text{-}4.3(\text{m}, 1\text{H}), 3.9\text{-}4.1 (\text{m}, 2\text{H}), 3.5\text{-}3.7(\text{m}, 2\text{H}), 1.7\text{-}2.6(\text{m}, 11\text{H}), 1.3\text{-}1.5(\text{m}, 1\text{H}), 1.03(\text{s}, 3\text{H})$

IR(KBr): $\nu(\text{cm}^{-1}) = 3422, 2934, 1688, 1636, 1580, 1539, 1483, 1431, 1304, 1233, 1186$.

40 Example 28

Preparation of 5-[3'-(4''-heptyloxy)phenoxybenzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 423 in Table 1):

45 The reaction was effected with 3-(4'-heptyloxy)phenoxybenzoic acid (296 mg, 1.26 mmol) in the same manner as in Example 7 to give the titled compound (297 mg, yield 56%).

m.p.153-156 °C

50 $^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.1\text{-}7.3(\text{m}, 4\text{H}), 7.00(\text{dd}, 1\text{H}, 1.4\text{Hz}, 8.0\text{Hz}), 5.98(\text{d}, 1\text{H}, 10\text{Hz}), 5.90(\text{t}, 1\text{H}, 3.4\text{Hz}), 4.31(\text{t}, 1\text{H}, 5.7\text{Hz}), 4.18(\text{ddd}, 1\text{H}, 5.2\text{Hz}, 10\text{Hz and } 10\text{Hz}), 2.53(\text{dd}, 1\text{H}, 5.6\text{Hz}, 18.7\text{Hz}), 2.1\text{-}2.5(\text{m}, 3\text{H}), 1.3\text{-}2.0(\text{m}, 12\text{H}), 1.01(\text{s}, 3\text{H}), 0.90(\text{t}, 6\text{H}, 7.2\text{Hz})$

IR(KBr): $\nu(\text{cm}^{-1}) = 3297, 2959, 2872, 1684, 1636, 1580, 1541, 1305, 1277, 1235$.

EP 0 579 223 A1

Example 29

Preparation of 5-[3'-(2'',4''-dimethyl-3''-pentyloxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 481 in Table 1):

5 The reaction was effected with 3-(2',4'-dimethyl-3'-pentyloxy)benzoic acid (235 mg, 0.99 mmol) in the same manner as in Example 7 to give the titled compound (203 mg, yield 51%).
m.p.190-192 °C

10 $^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.10\text{-}7.40(\text{m}, 4\text{H}), 7.04(\text{dd}, 1\text{H}, 2.7\text{Hz}, 8.1\text{Hz}), 5.97(\text{d}, 1\text{H}, 9.6\text{Hz}), 5.90(\text{dd}, 1\text{H}, \text{each } 3.7\text{Hz}), 4.10\text{-}4.15(\text{m}, 1\text{H}), 3.96(\text{t}, 1\text{H}, 5.8\text{Hz}), 2.1\text{-}2.6(\text{m}, 4\text{H}), 1.3\text{-}2.1(\text{m}, 6\text{H}), 1.01, 0.96, 0.93, 0.90\text{-}(\text{each s, each } 3\text{H}), 0.92(\text{s, 6H})$
IR(KBr): $\nu(\text{cm}^{-1}) = 3347, 2965, 1682, 1638, 1578, 1541, 1508, 1474, 1458, 1426, 1285, 1236, 1123$.

Example 30

15 Preparation of 5-(3'-dicyclopropylmethoxybenzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 529 in Table 1):

20 The reaction was effected with 3-dicyclopropylmethoxybenzoic acid (307 mg, 1.32 mmol) in the same manner as in Example 7 to give the titled compound (203 mg, yield 36%).
m.p.129-132 °C

25 $^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.23\text{-}7.40(\text{m}, 3\text{H}), 7.18(\text{br-s}, 1\text{H}), 7.03(\text{ddd}, 1\text{H}, 1.5\text{Hz}, 6.0\text{Hz}, 7.4\text{Hz}), 5.98\text{-}(\text{d}, 1\text{H}, 9.65\text{Hz}), 5.90\text{-}(\text{br-s}, 1\text{H}), 4.12\text{-}4.23(\text{m}, 1\text{H}), 3.53(\text{t}, 1\text{H}, 7.0\text{Hz}), 2.52(\text{dd}, 1\text{H}, 5.2\text{Hz}, 18\text{Hz}), 2.1\text{-}2.5(\text{m}, 3\text{H}), 1.7\text{-}1.95(\text{m}, 3\text{H}), 1.44(\text{ddd}, 1\text{H}, 5.5\text{Hz}, 5.5\text{Hz and } 13.1\text{Hz}), 1.0\text{-}1.2(\text{m}, 2\text{H}), 1.01(\text{s, 3H}), 0.6\text{-}0.8(\text{m, 4H}), 0.4\text{-}0.6(\text{m, 4H})$
IR(KBr): $\nu(\text{cm}^{-1}) = 3341, 3081, 3009, 2938, 1672, 1634, 1580, 1545, 1485, 1429, 1304, 1233, 1211, 1003, 982$.

Example 31

30 Preparation of 5-(3'-cyclohexylmethoxybenzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 424 in Table 1):

35 The reaction was effected with 3-cyclohexylmethoxybenzoic acid (472 mg, 2.01 mmol) in the same manner as in Example 7 to give the titled compound (406 mg, yield 48%).
m.p.159-161 °C

40 $^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.1\text{-}7.4(\text{m}, 4\text{H}), 7.01(\text{dd}, 1\text{H}, 1.3\text{Hz}, 7.0\text{Hz}), 5.99(\text{d}, 1\text{H}, 9.7\text{Hz}), 5.90(\text{dd}, 1\text{H}, \text{each } 3.9\text{Hz}), 4.1\text{-}4.3(\text{m}, 1\text{H}), 3.78(\text{d}, 2\text{H}, 6.1\text{Hz}), 2.1\text{-}2.6(\text{m}, 4\text{H}), 1.6\text{-}1.95(\text{m}, 9\text{H}), 1.45(\text{ddd}, 5.5\text{Hz}, 12.7\text{Hz and } 12.7\text{Hz}), 0.90\text{-}1.30(\text{m, 5H}), 1.01(\text{s, 3H})$
IR(KBr): $\nu(\text{cm}^{-1}) = 3275, 2930, 2853, 1680, 1636, 1611, 1582, 1543, 1449, 1429, 1319, 1277, 1240, 1038$.

Example 32

45 Preparation of 5-(3'-dicyclohexylmethoxybenzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 442 in Table 1):

50 The reaction was effected with 3-dicyclohexylmethoxybenzoic acid (293 mg, 0.93 mmol) in the same manner as in Example 7 to give the titled compound (257 mg, yield 54%).
m.p.142-146 °C

55 $^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.1\text{-}7.35(\text{m}, 4\text{H}), 7.03(\text{dd}, 1\text{H}, 1.8\text{Hz}, 8.0\text{Hz}), 5.98(\text{d}, 1\text{H}, 9.7\text{Hz}), 5.90(\text{dd}, 1\text{H}, \text{each } 3.6\text{Hz}), 4.1\text{-}4.3(\text{m}, 1\text{H}), 3.99(\text{t}, 1\text{H}, 5.8\text{Hz}), 2.1\text{-}2.6(\text{m}, 4\text{H}), 1.5\text{-}1.95(\text{m, 15H}), 1.4\text{-}1.5 (\text{m, 1H}), 1.0\text{-}1.3(\text{m, 10H}), 1.02(\text{s, 3H})$
IR(KBr): $\nu(\text{cm}^{-1}) = 3447, 2928, 2853, 1682, 1636, 1580, 1541, 1522, 1481, 1456, 1316, 1277, 1235$.

EP 0 579 223 A1

Example 33

Preparation of 5-[3'-(1"-morpholinomethyl)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 577 in Table 1):

The reaction was effected with 3-(1-morpholinomethyl)benzoic acid (500 mg, 2.26 mmol) in the same manner as in Example 1 to give the titled compound (327 mg, yield 35.1%).

m.p. (amorphous)

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.76(\text{s}, 1\text{H}), 7.64(\text{d}, 1\text{H}, J = 7.5\text{Hz}), 7.50(\text{d}, 1\text{H}, J = 7.5\text{Hz}), 7.40(\text{t}, 1\text{H}, J = 7.5\text{Hz}), 7.10(\text{d}, 1\text{H}, J = 1.9\text{Hz}), 5.87(\text{t}, 1\text{H}, J = 3.5\text{Hz}), 3.91-4.25(\text{m}, 5\text{H}), 3.70(\text{s}, 2\text{H}, 2.85-3.17 (\text{m}, 4\text{H}), 2.16-2.54(\text{m}, 4\text{H}), 1.81-1.92(\text{m}, 3\text{H}), 1.36-1.51(\text{m}, 1\text{H}), 1.12(\text{s}, 3\text{H})$

$\text{IR(KBr)}:\nu(\text{cm}^{-1}) = 3395, 2935, 1687, 1642, 1539, 1454, 1304, 1263, 1213, 1127, 1078.$

Example 34

Preparation of 5-[3'-(1"-piperidinomethyl)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 578 in Table 1):

The reaction was effected with 3-(1-piperidinomethyl)benzoic acid (338 mg, 1.54 mmol) in the same manner as in Example 1 to give the titled compound (328 mg, yield 52.2%).

m.p. (amorphous)

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 8.54(\text{s}, 1\text{H}), 7.96(\text{t}, 1\text{H}, J = 1.8\text{Hz}), 7.39, 7.51(\text{m}, 3\text{H}), 7.11(\text{d}, 1\text{H}, J = 1.9\text{Hz}), 5.82(\text{t}, 1\text{H}, J = 3.5\text{Hz}), 4.00-4.28(\text{m}, 3\text{H}), 2.84-3.22(\text{m}, 4\text{H}), 1.76-2.53(\text{m}, 9\text{H}), 1.54-1.71 (\text{m}, 2\text{H}), 1.35-1.50(\text{m}, 1\text{H}), 1.12(\text{s}, 3\text{H})$

$\text{IR(KBr)}:\nu(\text{cm}^{-1}) = 3422, 2942, 1644, 1541, 1454, 1316, 1213, 1046.$

Example 35

Preparation of 5-(3'-cyclohexylmethylbenzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 579 in Table 1):

The reaction was effected with 3-cyclohexylmethylbenzoic acid (352 mg, 1.61 mmol) in the same manner as in Example 7 to give the titled compound (296 mg, yield 63%).

m.p. 182-183 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.45-7.60(\text{m}, 2\text{H}), 7.15-7.40(\text{m}, 3\text{H}), 5.98(\text{d}, 1\text{H}, 12.2\text{Hz}), 5.90(\text{br-s}, 1\text{H}), 4.1-4.3(\text{m}, 1\text{H}), 2.52(\text{d}, 2\text{H}, 6.9\text{Hz}), 2.1-2.6(\text{m}, 4\text{H}), 1.35-2.00(\text{m}, 10\text{H}), 0.8-1.3(\text{m}, 5\text{H}), 1.02(\text{s}, 3\text{H})$

$\text{IR(KBr)}:\nu(\text{cm}^{-1}) = 3277, 2924, 2851, 1674, 1634, 1541, 1451, 1426, 1306, 1275.$

Example 36

Preparation of 5-(3'-cycloheptylmethylbenzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 581 in Table 1):

The reaction was effected with 3-cycloheptylmethyl benzoic acid (50.8 mg, 0.219 mmol) in the same manner as in Example 7 to give the titled compound (45.5mg, yield 49%).

m.p. 171-173 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.59(\text{br-s}, 1\text{H}), 7.52(\text{ddd}, 1\text{H}, 1.2\text{Hz}, 1.2\text{Hz and } 8.0\text{Hz}), 7.1-7.4(\text{m}, 3\text{H}), 5.98-\text{(d}, 1\text{H}, 8.9\text{Hz}), 5.92(\text{br-s}, 1\text{H}), 2.57(\text{d}, 2\text{H}, 7.2\text{Hz}), 2.1-2.6(\text{m}, 4\text{H}), 1.1-1.95(\text{m}, 17\text{H}), 1.02(\text{s}, 3\text{H})$

$\text{IR(KBr)}:\nu(\text{cm}^{-1}) = 3293, 2922, 2853, 1684, 1636, 1541, 1458, 1275, 1088.$

Example 37

Preparation of 5-[3-bromo-5-(p-isopropylphenoxy) benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 596 in Table 1):

The reaction was effected with 3-bromo-5-(p-isopropylphenoxy)benzoic acid (498 mg, 1.49 mmol) in the same manner as in Example 1 to give the titled compound (520 mg, yield 67%).

m.p. 62-65 °C

EP 0 579 223 A1

¹H-NMR(CDCl₃, 250MHz)δ = 7.51(dd, 1H, each 1.4Hz), 7.31(dd, 1H, each 1.4Hz), 7.15-7.25(m, 4H), 6.95-
5(d, 2H, 8.6Hz), 5.85-5.95(m, 2H), 4.05-4.20(m, 1H), 2.84-2.97(m, 1H), 2.1-2.6(m, 4H), 2.75-2.90 (m, 3H),
1.35-2.50(m, 1H), 1.23, 1.26(each s, each 3H)

IR(KBr):ν(cm⁻¹) = 3308, 3079, 2961, 1682, 1638, 1570, 1541, 1426, 1305, 1277, 1238, 1206, 1171, 858.

5

Example 38

Preparation of 5-[3',5'-di(p-isopropylphenoxy)benzoylamino]-10-methyl-Δ^{1(2),8(9)}-octalin-2-carboxylic acid
(Compound No. 597 in Table 1):

10

The reaction was effected with 3,5-di(p-isopropylphenoxy)benzoic acid (478 mg, 1.22 mmol) in the same manner as in Example 1 to give the titled compound (340 mg, yield 48%).

m.p.158-159 °C

¹H-NMR(CDCl₃, 250MHz)δ = 7.10-7.20(m, 5H), 7.02(d, 2H, 2.2Hz), 6.94(d, 4H, 8.6Hz), 6.71(dd, 1H, each 2.2Hz), 5.82-5.90(m, 2H), 4.02 -4.20(m, 1H), 2.80-2.95(m, 2H), 2.10-2.60(m, 4H), 2.70-2.90(m, 3H), 1.30-1.50(m, 3H), 1.30-1.50(m, 1H), 1.21, 1.24(each s, each 3H), 0.95(s, 3H)

IR(KBr):ν(cm⁻¹) = 3291, 2963, 2936, 2872, 1682, 1640, 1609, 1588, 1555, 1505, 1435, 1327, 1283, 1217, 1173, 1123, 1005, 835.

20

Example 39

Preparation of 5-(3'-cycloheptyloxy-4'-methoxybenzoylamino)-10-methyl-Δ^{1(2),8(9)}-octalin-2-carboxylic acid
(Compound No. 139 in Table 1):

25

The reaction was effected with 3-cycloheptyloxy-4-methoxybenzoic acid (319 mg, 1.21 mmol) in the same manner as in Example 7 to give the titled compound (278 mg, yield 50%).

m.p.127-129 °C

¹H-NMR(CDCl₃, 250MHz)δ = 7.38(d, 1H, 2.0Hz), 7.17-7.21(m, 2H), 6.85(d, 1H, 8.5Hz), 5.89-5.95(m, 2H), 4.39-4.50(m, 1H), 4.11-4.23 (m, 1H), 3.87(s, 3H), 2.52(dd, 1H, 5.0Hz, 18.1Hz), 2.01-2.48(m, 5H), 1.35-1.95(m, 14H), 1.01(s, 3H)

IR(KBr):ν(cm⁻¹) = 3447, 3291, 2932, 2861, 1680, 1634, 1507, 1269.

Example 40

35

Preparation of 5-(3',4'-methylenedioxybenzoylamino)-10-methyl-Δ^{1(2),8(9)}-octalin-2-carboxylic acid
(Compound No. 491 in Table 1):

The reaction was effected with piperonylic acid (217 mg, 1.31 mmol) in the same manner as in Example 7 to give the titled compound (293 mg, yield 61%).

40

m.p.(amorphous)

¹H-NMR(CDCl₃, 250MHz)δ = 7.20-7.30(m, 2H), 7.18(d, 1H, 1.7Hz), 6.82(d, 1H, 7.4Hz), 6.01(s, 2H), 5.91-5.95(m, 2H), 4.10-4.21(m, 1H), 4.52(dd, 1H, 4.1Hz, 18Hz), 2.1-2.5(m, 3H), 1.75-1.95(m, 3H), 1.42(ddd, 1H, 5.5Hz, 12.7Hz and 12.7Hz), 1.00(s, 3H)

IR(KBr):ν(cm⁻¹) = 3314, 2934, 1699, 1636, 1541, 1505, 1487, 1439, 1259, 1209, 1182, 1038.

45

Example 41

Preparation of 5-(3',4'-ethylenedioxybenzoylamino)-10-methyl-Δ^{1(2),8(9)}-octalin-2-carboxylic acid (Compound No. 493 in Table 1):

50

The reaction was effected with 3,4-ethylenedioxybenzoic acid (214 mg, 1.19 mmol) in the same manner as in Example 7 to give the titled compound (188 mg, yield 43%).

m.p.(amorphous)

¹H-NMR(CDCl₃, 250MHz)δ = 7.27-7.30(m, 2H), 7.17(br-s, 1H), 6.89(d, 1H, 8.3Hz), 5.84-5.90(m, 2H), 4.27-

55

(s, 4H), 4.08-4.25 (m, 1H), 2.51(dd, 1H, 4.5Hz, 18.4Hz), 2.10-2.47(m, 3H), 1.75-1.90 (m, 3H), 1.42(ddd, 1H, 5.7Hz, 5.7Hz, 13.1Hz), 1.00(s, 3H)

IR(KBr):ν(cm⁻¹) = 3341, 2936, 1684, 1634, 1615, 1582, 1541, 1501, 1316, 1289, 1260, 1067.

Example 42

Preparation of 5-(3',4'-cyclohexylidenedioxybenzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 495 in Table 1):

The reaction was effected with 3,4-cyclohexylidenedioxybenzoic acid (303 mg, 1.29 mmol) in the same manner as in Example 7 to give the titled compound (250 mg, yield 46%).

m.p.104-105 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.24(\text{dd}, 1\text{H}, 1.8\text{Hz}, 8.1\text{Hz}), 7.14\text{-}7.18 (\text{m}, 2\text{H}), 6.73(\text{d}, 1\text{H}, 8.1\text{Hz}), 5.8\text{-}5.95\text{ (m, }2\text{H}), 4.15(\text{ddd}, 1\text{H}, 4.9\text{Hz, }10\text{Hz and }10\text{Hz}), 2.51(\text{dd}, 1\text{H}, 5.3\text{Hz, }19\text{Hz}), 2.15\text{-}2.45(\text{m, }3\text{H}), 1.82\text{-}1.95 (\text{m, }7\text{H}), 1.35\text{-}1.60(\text{m, }5\text{H}), 0.99(\text{s, }3\text{H})$

$\text{IR(KBr)}:\nu(\text{cm}^{-1}) = 3301, 2940, 1866, 1674, 1636, 1559, 1541, 1491, 1437, 1360, 1306, 1283, 1258.$

Example 43

Preparation of 5-(3',5'-di-t-butylbenzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 503 in Table 1):

The reaction was effected with 3,5-di-t-butylbenzoic acid (243 mg, 1.04 mmol) in the same manner as in Example 7 to give the titled compound (143 mg, yield 32%).

m.p.99-101 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.55\text{-}7.57(\text{m, }3\text{H}), 7.18(\text{br-s, }1\text{H}), 5.90\text{-}5.96(\text{m, }2\text{H}), 4.16\text{-}4.30(\text{m, }1\text{H}), 2.53\text{ (dd, }1\text{H, }5.0\text{Hz, }19\text{Hz}), 2.10\text{-}2.50(\text{m, }3\text{H}), 1.70\text{-}1.90(\text{m, }3\text{H}), 1.46(\text{ddd, }1\text{H, }5.7\text{Hz, }5.7\text{Hz, }13\text{Hz}), 1.34(\text{s, }18\text{H}), 1.02(\text{s, }3\text{H})$

$\text{IR(KBr)}:\nu(\text{cm}^{-1}) = 3293, 2961, 1682, 1634, 1595, 1539, 1474, 1458, 1424, 1364, 1265, 1213, 706.$

Example 44

Preparation of 5-(5',6',7',8'-tetrahydro-5',5',8',8'-tetramethylnaphthalene-2'-carbonylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 509 in Table 1):

The reaction was effected with 5,6,7,8-tetrahydro-5,5,8,8-tetramethylnaphthalene-2-carboxylic acid (207 mg, 0.89 mmol) in the same manner as in Example 7 to give the titled compound (232 mg, yield 62%).

m.p.109-110 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.80(\text{d, }1\text{H, }1.8\text{Hz}), 7.41(\text{dd, }1\text{H, }1.9\text{Hz, }8.2\text{Hz}), 7.33(\text{d, }1\text{H, }8.2\text{Hz}), 7.17(\text{d, }1\text{H, }1.5\text{Hz}), 5.95(\text{d, }1\text{H, }9.7\text{Hz}), 5.90(\text{dd, }1\text{H, each }3.7\text{Hz}), 4.10\text{-}4.25(\text{m, }1\text{H}), 2.10\text{-}2.60(\text{m, }4\text{H}), 1.78\text{-}1.95(\text{m, }3\text{H}), 1.68(\text{s, }4\text{H}), 1.44(\text{ddd, }1\text{H, }5.5\text{Hz, }12.7\text{Hz and }12.7\text{Hz}), 1.29, 1.30(\text{each s, each }3\text{H}), 1.27(\text{s, }6\text{H}), 1.01(\text{s, }3\text{H})$

$\text{IR(KBr)}:\nu(\text{cm}^{-1}) = 3310, 2961, 1684, 1634, 1559, 1539, 1267.$

Example 45

Preparation of (5R,10R)-5-(3'-cycloheptyloxybenzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 111 in Table 1):

The reaction was effected with 3-cycloheptyloxybenzoic acid (359 mg, 1.53 mmol) and (5R,10R)-5-amino-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid methyl ester (339 mg, 1.53 mmol) in the same manner as in Example 7 to give the titled compound (370 mg, yield 57%).

$[\alpha]_D^{26} = +135^\circ (c = 0.50, \text{MeOH})$

m.p.133-135 °C

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.1\text{-}7.4(\text{m, }4\text{H}), 6.98\text{-}7.02(\text{m, }1\text{H}), 6.01(\text{br-d, }1\text{H, }9.7\text{Hz}), 5.92(\text{t-like, }1\text{H, }J=1.0\text{Hz}), 4.20\text{-}4.35(\text{m, }1\text{H}), 4.1\text{-}4.3(\text{m, }1\text{H}), 2.1\text{-}2.6(\text{m, }4\text{H}), 2.0\text{-}2.1(\text{m, }2\text{H}), 1.3\text{-}1.9(\text{m, }8\text{H}), 1.03(\text{s, }3\text{H})$

Example 46

Preparation of (5S,10S)-5-(3'-cycloheptyloxybenzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 111 in Table 1):

The reaction was effected with 3-cycloheptyloxybenzoic acid (280 mg, 1.19 mmol) and (5S,10S)-5-amino-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid methyl ester (264 mg, 1.19 mmol) in the same manner as in Example 7 to give the titled compound (271 mg, yield 54%).

$$[\alpha]_D^{26} = -128^\circ \text{ (c = 0.50, MeOH)}$$

¹H-NMR(CDCl₃, 250MHz) δ = 7.1-7.4(m, 4H), 6.98-7.02(m, 1H), 6.01(br-d, 1H, 9.7Hz), 5.92(t-like, 1H, J=1.0Hz), 4.20-4.35 (m, 1H), 4.1-4.3(m, 1H), 2.1-2.6(m, 4H), 2.0-2.1(m, 2H), 1.3-1.9 (m, 8H), 1.03(s, 3H).

Example 47

Preparation of (5S,10S)-5-(3'-dicyclopropylmethoxybenzoylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 529 in Table 1):

The reaction was effected with 3-dicyclopropylmethoxybenzoic acid (105 mg, 0.48 mmol) and (5S,10S)-5-amino-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid methyl ester (110 mg, 0.48 mmol) in the same manner as in Example 7 to give the titled compound (56 mg, yield 27%).

$$[\alpha]_D^{27} = -126^\circ \text{ (c = 0.50, MeOH)}$$

m.p. 97-99 °C

¹H-NMR(CDCl₃, 250MHz) δ = 7.23-7.40(m, 3H), 7.18(br-s, 1H), 7.03(ddd, 1H, 1.5Hz, 6.0Hz, 7.4Hz), 5.98(d, 1H, 9.65Hz), 5.90(br-s, 1H), 4.12-4.23(m, 1H), 3.53(t, 1H, 7.0Hz), 2.52(dd, 1H, 5.2Hz, 18Hz), 2.1-2.5(m, 3H), 1.7-1.95(m, 3H), 1.44(ddd, 1H, 5.5Hz, 5.5Hz and 13.1Hz), 1.0-1.2(m, 2H), 1.01(s, 3H), 0.6-0.8(m, 4H), 0.4-0.6(m, 4H).

Example 48

Preparation of (5S,10S)-5-[3'-(2'',4''-dimethyl-3''-pentyloxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 481 in Table 1):

The reaction was effected with 3-(2',4'-dimethyl-3'-pentyloxy)benzoic acid (124 mg, 0.52 mmol) and (5S,10S)-5-amino-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid methyl ester (116 mg, 0.52 mmol) in the same manner as in Example 7 to give the titled compound (216 mg, yield 66%).

$$[\alpha]_D^{27} = -170^\circ \text{ (c = 1.0, MeOH)}$$

m.p. 116-118 °C

¹H-NMR(CDCl₃, 250MHz) δ = 7.10-7.40(m, 4H), 7.04(dd, 1H, 2.7Hz, 8.1Hz), 5.97(d, 1H, 9.6Hz), 5.90(dd, 1H, each 3.7Hz), 4.10-4.15(m, 1H), 3.96(t, 1H, 5.8Hz), 2.1-2.6(m, 4H), 1.3-2.1(m, 6H), 1.01, 0.96, 0.93, 0.90-(each s, each 3H), 0.92(s, 6H)

IR(KBr): ν (cm⁻¹) = 3328, 2965, 2874, 1682, 1634, 1580, 1532, 1481, 1429, 1316, 1275, 1236, 1213, 1186, 1005, 752.

Example 49

Preparation of (5S,10S)-N-methyl-5-[3'-(2'',4''-dimethyl-3''-pentyloxy)benzoylamino]-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 497 in Table 1):

The reaction was effected with 3-(2',4'-dimethyl-3'-pentyloxy)benzoic acid (137 mg, 0.58 mmol) and (5S,10S)-5-amino-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid methyl ester (128 mg, 0.58 mmol) in the same manner as in Example 7 to give the titled compound (238 mg, yield 99%). This compound was dissolved in dimethylformamide (1 ml) and mixed with sodium hydride (27 mg, 0.66 mmol), 0.5 hours later mixed with methyl iodide (0.055 ml, 0.89 mmol), and the resultant mixture was stirred for 3 hours. The reaction mixture was poured into chilled water, extracted with diethyl ether, and the extract was concentrated. The product was hydrolyzed and purified in the same manner as in Example 7 to give the titled compound (170 mg, yield 70%).

$$[\alpha]_D^{27} = -166^\circ \text{ (c = 0.33, MeOH)}$$

m.p.(amorphous)

EP 0 579 223 A1

¹H-NMR(CDCl₃, DMSO-d₆, 373K, 250MHz)δ = 7.27(dd, 1H, each 7.9Hz), 6.96(dd, 1H, 1.8Hz, 7.9Hz), 6.80-6.90(m, 3H), 5.76(br-s, 1H), 5.76(br-s, 1H), 3.99(t, 1H, 5.6Hz), 2.90(s, 3H), 2.1-2.5(m, 5H), 1.9-2.1(m, 2H), 1.65-1.85(m, 2H), 1.1-1.3(m, 1H), 1.05(s, 3H), 0.9-1.0(m, 12H)

IR(KBr):ν(cm⁻¹) = 2965, 2874, 1701, 1684, 1634, 1578, 1456, 1404, 1364, 1316, 1256, 1209, 1005, 980, 793, 754.

Example 50

Preparation of 5-(3'-cycloheptyloxy-4'-isopropoxybenzoylamino)-10-methyl-Δ^{1(2),8(9)}-octalin-2-carboxylic acid (Compound No. 117 in Table 1):

The reaction was effected with 3-cycloheptyloxy-4-isopropoxybenzoic acid (157 mg, 0.54 mmol) and 5-amino-10-methyl-Δ^{1(2),8(9)}-octalin-2-carboxylic acid methyl ester (125 mg, 0.54 mmol) in the same manner as in Example 7 to give the titled compound (128 mg, yield 48%).

m.p.143-196 °C

¹H-NMR(CDCl₃, 250MHz)δ = 7.38(d, 1H, 2.1Hz), 7.23(dd, 1H, 2.1Hz, 8.3Hz), 7.19(br-s, 1H), 6.89(d, 1H, 8.3Hz), 5.90-5.95 (m, 2H), 4.49-4.62(m, 1H), 4.30-4.46(m, 1H), 4.12-4.23(m, 1H), 2.53(dd, 1H, 5.1Hz, 19Hz), 2.15-2.48(m, 3H), 1.3-2.1(m, 16H), 1.32, 1.34(each s, each 3H), 1.01(s, 3H)

IR(KBr):ν(cm⁻¹) = 2976, 2932, 2859, 1682, 1634, 1601, 1499, 1302, 1217, 1138, 1107, 1001, 954.

Example 51

Preparation of 5-(3'-cycloheptyloxy-2-methylbenzoylamino)-10-methyl-Δ^{1(2),8(9)}-octalin-2-carboxylic acid (Compound No. 119 in Table 1):

The reaction was effected with 3-cycloheptyloxy-2-methylbenzoic acid (200 mg, 0.81 mmol) in the same manner as in Example 1 to give the titled compound (297 mg, yield 84.1%).

m.p.237-239 °C

¹H-NMR(CDCl₃, 250MHz)δ = 7.20(s, 1H), 7.16(dd, 1H, J = 7.2Hz, 8.3Hz), 6.91(d, 1H, J = 7.2Hz), 6.86(d, 1H, J = 8.3Hz), 5.92(t, 1H, J = 3.7Hz), 5.64(d, 1H, J = 9.7Hz), 4.40-4.49(m, 1H), 4.14-4.24(m, 1H), 2.16-2.66(m, 3H), 2.29(s, 3H), 1.40-2.07(m, 17H), 0.96(s, 3H)

IR(KBr):ν(cm⁻¹) = 3277, 2932, 2858, 1672, 1634, 1541, 1458, 1425, 1308, 1260, 1186, 1078.

Example 52

Preparation of (1S,8S)-1-(3'-cycloheptyloxybenzoylamino)-8a-methyl-1,2,6,7-tetrahydroindene-5-carboxylic acid (Compound No. 135 in Table 1):

The reaction was effected with 3-cycloheptyloxybenzoic acid (104 mg, 0.44 mmol) and (1S,8S)-1-amino-8a-methyl-1,2,6,7-tetrahydroindene-5-carboxylic acid methyl ester (92 mg, 0.44 mmol) in the same manner as in Example 7 to give the titled compound (18 mg, yield 9.3%).

¹H-NMR(CDCl₃, 250MHz)δ = 7.1-7.3(m, 4H), 6.89(dd, 1H, 1.8Hz, 7.4Hz), 6.73(br-d, 1H, 8.7Hz), 6.04(br-s, 1H), 4.3-4.5(m, 2H), 2.63(ddd, 1H, 3Hz, 7.6Hz, 17Hz), 2.1-2.5(m, 3H), 1.85-2.0(m, 3H), 1.30-1.8(m, 2H), 0.92-1.0(s, 1H)

IR(KBr):ν(cm⁻¹) = 3380, 2930, 2857, 1686, 1640, 1580, 1483, 1238, 1181, 1086, 1019, 696.

Example 53

Preparation of (1S,8S)-1-[3'-(2'',4''-dimethyl-3''-pentyl)oxybenzoylamino]-8a-methyl-1,2,6,7-tetrahydroindene-5-carboxylic acid (Compound No. 505 in Table 1):

The reaction was effected with 3-(2',4'-dimethyl-3'-pentyl)oxybenzoic acid (64 mg, 0.27 mmol) and 1-amino-8a-methyl-1,2,6,7-tetrahydroindene-5-carboxylic acid methyl ester (56 mg, 0.27 mmol) in the same manner as in Example 7 to give the titled compound (25 mg, yield 22%).

¹H-NMR(CDCl₃, 250MHz)δ = 7.1-7.4(m, 4H), 7.05(dd, 1H, 1.8Hz, 8.1Hz), 6.21(d, 1H, 8.8Hz), 5.87(br-s, 1H), 4.45-4.6(m, 1H), 3.96 (t, 1H, 5.7Hz), 2.81(ddd, 1H, 3.0Hz, 7.6Hz, 16.8Hz), 2.3-2.7(m, 3H), 1.9-2.1(m, 3H), 1.59(ddd, 5.7Hz, 5.7Hz and 12.5Hz), 0.96, 0.95, 0.93, 0.92, 0.90(each s, each 3H)

IR(KBr):ν(cm⁻¹) = 3374, 2965, 1686, 1638, 1580, 1539, 1481, 1283, 1240, 1198, 1003, 754.

Example 54

Preparation of (1S,8S)-1-[3'-(dicyclopropylmethoxy)benzoylamino]-8a-methyl-1,2,6,7-tetrahydroindene-5-carboxylic acid (Compound No. 553 in Table 1):

The reaction was effected with 3-(dicyclopropylmethoxy)benzoic acid (72 mg, 0.31 mmol) and 1-amino-8a-methyl-1,2,6,7-tetrahydroindene-5-carboxylic acid methyl ester (64 mg, 0.31 mmol) in the same manner as in Example 7 to give the titled compound (29 mg, yield 22%).

¹H-NMR(CDCl₃, 250MHz) δ = 7.2-7.4(m, 4H), 7.05(ddd, 1H, 2.2Hz, 2.2Hz, and 6.9Hz), 6.21(d, 1H, 8.9Hz), 5.87(br-s, 1H), 4.47-4.59 (m, 1H), 3.53(t, 1H, 7.0Hz), 2.81(ddd, 1H, 3Hz, 7.8Hz, 17Hz), 2.2-2.7(m, 3H), 2.04(dd, 1H, 4.2Hz, 13Hz), 1.59(ddd, 5.7Hz, 5.7Hz, and 12.5Hz), 1.0-1.2(m, 2H), 1.03(s, 3H), 0.4-0.6(m, 4H), 0.2-0.4(m, 4H)

IR(KBr): ν (cm⁻¹) = 3364, 3083, 3009, 2961, 2930, 1686, 1638, 1582, 1539, 1483, 1296, 1240, 1198.

Example 55

Preparation of 5-(3'-cycloheptyloxybenzenesulfonylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 620 in Table 1):

To a mixture of 5-amino-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid methyl ester (108 mg, 0.49 mmol), triethylamine (0.65 ml) and methylene chloride (2.6 ml) was added 3-cycloheptyloxybenzenesulfonyl chloride (129 mg, 0.47 mmol), and the resultant mixture was stirred at room temperature for 1 hour, poured into chilled water and extracted with ethyl acetate. The extract was concentrated, and the residue was chromatographed on a silica gel column to give crude product (88 mg). This product was hydrolyzed in the same manner as in Example 7 to give the titled compound (62 mg, yield 30%).

m.p.124-125 °C

¹H-NMR(CDCl₃, 250MHz) δ = 7.30-7.40(m, 3H), 7.09(br-s, 1H), 7.03(ddd, 1H, 2.4Hz and 6.8Hz), 5.77(t, 1H, 3.7Hz), 4.4-4.5(m, 1H), 4.38(br-d, 1H, 9.8Hz), 3.05-3.20(m, 1H), 2.45(dd, 1H, 5.1Hz, 18.4Hz), 2.1-2.3(m, 3H), 1.3-2.1(m, 15H), 0.95-1.10(m, 1H), 0.87(s, 3H)

IR(KBr): ν (cm⁻¹) = 3268, 2932, 2863, 2631, 2540, 1682, 1632, 1613, 1476, 1431, 1323, 1289, 1252, 1236, 1157, 1096, 1063.

Example 56

Preparation of (5S,10S)-5-(3'-cycloheptyloxybenzene-sulfonylamino)-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid (Compound No. 620 in Table 1):

The reaction was effected with 3-cycloheptyloxy benzenesulfonyl chloride (142 mg, 0.51 mmol) and (5S,10S)-5-amino-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid methyl ester (87 mg, 0.40 mmol) in the same manner as in Example 55 to give the titled compound (82 mg, yield 46%).

m.p.70-72 °C

¹H-NMR(CDCl₃, 250MHz) δ = 7.30-7.40(m, 3H), 7.09(br-s, 1H), 7.03(ddd, 1H, 2.4Hz and 6.8Hz), 5.77(t, 1H, 3.7Hz), 4.4-4.5(m, 1H), 4.38(br-d, 1H, 9.8Hz), 3.05-3.20(m, 1H), 2.45(dd, 1H, 5.1Hz, 18.4Hz), 2.1-2.3(m, 3H), 1.3-2.1(m, 15H), 0.95-1.10(m, 1H), 0.87(s, 3H).

Example 57

Preparation of 5-(3'-phenoxybenzoylamino)-5,6,7,8-tetrahydro-naphthalene-2-carboxylic acid (Compound No. 624 in Table 2):

The reaction was effected with 3-phenoxybenzoic acid (0.91 mg, 4.23 mmol) and 1-amino-6-trifluoromethanesulfonyl-1,2,3,4-tetrahydronaphthalene (1.5 g, 5.08 mmol) to give the condensation product (1.5 g, 73%). This product (0.537 mg, 1.09 mmol) was added to a mixture of palladium acetate (123 mg, 0.55 mmol), triphenyl phosphine (286 mg, 1.1 mmol), triethylamine (19.4 ml), methanol (50 ml) and tetrahydrofuran (50 ml), and the resultant mixture was reacted in an atmosphere of carbon monoxide for 24 hours. The reaction mixture was poured into water and extracted with ethyl acetate. The extract was concentrated, and the residue was hydrolyzed in the same manner as in Example 7 to give the titled compound (88 mg, yield 21%).

m.p.155-160 °C

$^1\text{H-NMR}(\text{CDCl}_3 + \text{catCD}_3\text{OD}, 250\text{MHz})\delta = 7.81(\text{d-like}, 2\text{H}, 4\text{Hz}), 7.30\text{-}7.55(\text{m}, 5\text{H}), 7.10\text{-}7.20(\text{m}, 2\text{H}), 6.95\text{-}7.10(\text{m}, 2\text{H}), 6.62\text{-}6.78 (\text{m}, 1\text{H}), 5.30\text{-}5.45(\text{m}, 1\text{H}), 2.80\text{-}2.95(\text{m}, 2\text{H}), 2.1\text{-}2.3(\text{m}, 1\text{H}), 2.85\text{-}2.95(\text{m}, 3\text{H})$

$\text{IR(KBr)}:\nu(\text{cm}^{-1}) = 3274, 2936, 1694, 1632, 1578, 1534, 1481, 1429, 1269, 1233, 1200, 752, 691.$

5

Example 58

Preparation of 5-(3'-phenoxybenzoyl)amino-9-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-(2'-trans-propenoic acid) (Compound No. 640 in Table 3):

10

To a mixture of 3-phenoxybenzoic acid (93 mg, 0.44 mmol), triethylamine (0.31 ml, 2.2 mmol) and methylene chloride (3 ml) chilled at 0 °C was gradually added thionyl chloride (48 μl , 0.66 mmol), and the resultant mixture was stirred at 0 °C for 15 minutes. A solution of 5-amino-9-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-(2'-propenoic acid) methyl ester (114 mg, 0.44 mmol) in methylene chloride (3 ml) was added to the mixture, which was stirred at room temperature for 30 minutes. The reaction mixture was distributed between ethyl acetate and water, and the organic layer was washed with saturated brine and concentrated. The residue was dissolved in methanol (5 ml), mixed with 2 N aqueous KOH (1 ml) and refluxed under heating for 1 hour. The reaction mixture was acidified with 2 N hydrochloric acid, extracted with chloroform, dried and concentrated. The resulting residue was chromatographed on a silica gel column, eluting with chloroform-methanol to give the titled compound (60 mg, yield 32%).

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.34\text{-}7.49(\text{m}, 6\text{H}), 7.15(\text{t}, 2\text{H}, J = 7.5\text{Hz}), 7.03(\text{d}, 2\text{H}, J = 7.5\text{Hz}), 6.36(\text{br s}, 1\text{H}), 5.98(\text{d}, 1\text{H}, J = 9.7\text{Hz}), 5.85(\text{d}, 1\text{H}, J = 15.7\text{Hz}), 5.77(\text{t}, 1\text{H}, J = 3.7\text{Hz}), 4.16\text{-}4.31(\text{m}, 1\text{H}), 2.15\text{-}2.59(\text{m}, 3\text{H}), 1.82\text{-}1.92(\text{m}, 4\text{H}), 1.41\text{-}1.54(\text{m}, 1\text{H}), 1.03(\text{s}, 3\text{H})$

$\text{IR(KBr)}:\nu(\text{cm}^{-1}) = 3426, 3289, 3065, 2938, 1674, 1634, 1605, 1580, 1547, 1481, 1318, 1277, 1232.$

25

Example 59

Preparation of 1-[5'-(3''-phenoxybenzoylamino)-5',6',7',8'-tetrahydro-2'-naphthoxy]acetic acid (Compound No. 656 in Table 4):

30

The reaction was effected with 3-phenoxybenzoic acid (436 mg, 2.04 mmol) and 1-(5-amino-5',6',7',8'-tetrahydro-2'-naphthoxy)acetic acid methyl ester (479 mg, 2.04 mmol) in the same manner as in Example 1 to give the titled compound (300 mg, yield 35%).

m.p.153-1 57 °C

35

$^1\text{H-NMR}(\text{CDCl}_3 + \text{catDMSO-d}_6, 250\text{MHz})\delta = 7.35\text{-}7.45(\text{m}, 1\text{H}), 6.80\text{-}7.25(\text{m}, 9\text{H}), 6.58(\text{dd}, 1\text{H}, 2.7\text{Hz}, 8.5\text{Hz}), 5.1\text{-}5.2(\text{m}, 1\text{H}), 4.41(\text{s}, 2\text{H}), 2.5\text{-}2.7(\text{m}, 2\text{H}), 1.5\text{-}2.0(\text{m}, 4\text{H})$

$\text{IR(KBr)}:\nu(\text{cm}^{-1}) = 3343, 2932, 2866, 1725, 1611, 1574, 1549, 1481, 1435, 1333, 1289, 1186, 1163, 1132, 1080, 760, 694.$

40

Example 60

Preparation of 1-[5'-(3''-benzyloxybenzoylamino)-5',6',7',8'-tetrahydro-2'-naphthoxy]acetic acid (Compound No. 666 in Table 4):

45

The reaction was effected with 3-benzyloxybenzoic acid (398 mg, 1.74 mmol) and 1-(5'-amino-5',6',7',8'-tetrahydro-2'-naphthoxy)acetic acid methyl ester (410 mg, 1.74 mmol) in the same manner as in Example 1 to give the titled compound (127 mg, yield 17%).

m.p.(amorphous)

$^1\text{H-NMR}(\text{CDCl}_3 + \text{catDMSO-d}_6, 250\text{MHz})\delta = 7.1\text{-}7.4(\text{m}, 8\text{H}), 6.95\text{-}7.05 (\text{m}, 1\text{H}), 6.5\text{-}6.8(\text{m}, 3\text{H}), 5.15\text{-}5.25(\text{m}, 1\text{H}), 5.01(\text{s}, 2\text{H}), 4.47(\text{s}, 2\text{H}), 2.6\text{-}2.8(\text{m}, 2\text{H}), 1.95\text{-}2.10(\text{m}, 1\text{H}), 1.7\text{-}1.9(\text{m}, 3\text{H})$

50

$\text{IR(KBr)}:\nu(\text{cm}^{-1}) = 3345, 2928, 2859, 1723, 1613, 1578, 1551, 1499, 1445, 1290, 1240, 1217, 1163, 1132, 1080, 1017, 756, 698.$

55

Reference Example 1

Preparation of 5-amino-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid methyl ester:

To a mixture of Wieland Mischer's ketone (38.5 g, 216 mmol), diisopropylethylamine (39.5 ml) and methylene chloride (770 ml) was added anhydrous trifluoromethanesulfonic acid (40 ml, 238 mmol) under ice cooling, and the resultant mixture was stirred at room temperature for 2 hours. The reaction mixture was mixed with hexane (120 ml), and the organic layer was washed with water, 3 N hydrochloric acid, saturated aqueous sodium bicarbonate solution and saturated brine in this order, dried and concentrated to give monoenol triflate compound (63 g). This product was dissolved in a mixture of methanol (630 ml) and tetrahydrofuran (630 ml), mixed with palladium acetate (1.14 g, 5.08 mmol), triphenyl phosphine (2.6 g, 10.1 mmol) and triethylamine (315 ml). Carbon monoxide was bubbled into the solution at room temperature for 50 hours. Then, the reaction mixture was diluted with hexane, poured into chilled water and extracted with ethyl acetate to give light yellow syrup (33 g). This product was dissolved in methanol (1.3 L), mixed with ammonium acetate (115 g, 1.45 mmol) and sodium cyanoborohydride (6.4 g, 102 mmol), and the mixture was refluxed for 6 hours. The reaction mixture was allowed to warm at room temperature, poured into chilled water and extracted with chloroform under weakly basic conditions. The organic layer was dried and concentrated. The residue was washed with hexane-diethyl ether to give the titled compound (16.2 g, yield 34%) as white powders.

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.05(\text{d}, 1\text{H}), 5.78(\text{t}, 1\text{H}), 3.73(\text{s}, 3\text{H}), 2.62(\text{d}, 1\text{H}), 2.57(\text{dd}, 1\text{H}), 2.2-2.4(\text{m}, 3\text{H}), 1.95(\text{ddd-like}, 1\text{H}), 1.5-1.8(\text{m}, 3\text{H}), 1.1-1.3(\text{m}, 2\text{H}), 0.86(\text{s}, 3\text{H}).$

Reference Example 2

Preparation of (5S,10S)-5-amino-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid methyl ester:

The reaction was effected with (+)-Wieland Mischer's ketone (Fulka Company, $[\alpha]_D > +98^\circ$ (c = 1.0, benzene) (2.0 g, 11.2 mmol) in the same manner as in Reference Example 1 to give the titled compound (0.95 g, yield 57%).

$[\alpha]_D^{27} = 254^\circ$ (c = 0.50, MeOH)

Reference Example 3

Preparation of (5R,10R)-5-amino-10-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-carboxylic acid methyl ester:

The racemate obtained in Reference Example 1 (2.27 g, 10.3 mmol) was recrystallized three times together with L-benzoyltartaric acid (3.86 g, 10.3 ml) in methanol. The crystals are collected under basic conditions and purified to give the titled compound (339 mg, yield 15%).

$[\alpha]_D^{26} = +244^\circ$ (c = 0.5, MeOH)

Reference Example 4

Preparation of (1S,8S)-1-amino-8a-methyl-1,2,6,7-tetrahydro-indane-5-carboxylic acid methyl ester:

The reaction was effected with (+)-2,3,6,7-tetrahydro-8a-methyl-1,5-indadione ($[\alpha]_D^{29} > +358^\circ$ (c = 1.0, benzene) (2.95 g, 18.0 mmol) in the same manner as in Reference Example 1 to give the titled compound (408 g, yield 11%).

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.19(\text{d}, 1\text{H}, 2.4\text{Hz}), 5.74(\text{br-s-like}), 3.72(\text{s}, 3\text{H}), 3.10(\text{dd}, 1\text{H}, 7.6\text{Hz}, 10.0\text{Hz}), 1.8-2.7(\text{m}, 4\text{H}), 1.2-1.4 (\text{m}, 2\text{H}), 0.81(\text{s}, 3\text{H}).$

Reference Example 5

Preparation of 1-(1'-amino-1',2',3',4'-tetrahydro-2'-naphthoxy)acetic acid methyl ester:

The reductive amination was effected with 1-(1'-oxo-1',2',3',4'-tetrahydro-2'-naphthoxy)acetic acid methyl ester (9.0 g, 36.2 mmol), prepared from α -bromoacetic acid and 6-hydroxytetralone, in the same manner as in Reference Example 1 to give the titled compound (4.08 g, yield 48%).

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.13(\text{H}, \text{d}, 8.6\text{Hz}), 6.67(\text{H}, \text{dd}, 2.7\text{Hz and } 8.6\text{Hz}), 6.56(\text{d}, 1\text{H}, 2.7\text{Hz}), 4.97-$

EP 0 579 223 A1

5.10(m, 1H), 4.55(s, 2H), 3.75 (s, 3H), 2.60-2.80(m, 2H), 1.65-2.05(m, 4H).

Reference Example 6

5 Preparation of 5-amino-9-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-(2'-propenyllic acid) ethyl ester:

To a solution of palladium acetate (126 mg, 0.56 mmol), triphenyl phosphine (292 mg, 1.12 mmol) and tetrahydrofuran (2.9 ml) previously prepared was dropwise added a solution of enol triflate (1.74 g, 5.61 mmol) as disclosed in Reference Example 1 in tetrahydrofuran (10 ml). The mixture was mixed with 10 triethylamine (10 ml) and ethyl acetate (1.22 ml, 11.2 mmol), and stirred at room temperature for 3 days. The reaction mixture was diluted with ethyl acetate, and the organic layer was washed with water and saturated sodium bicarbonate solution, dried and concentrated. The residue was chromatographed on a silica gel column (40 g) eluting with hexane-ethyl acetate to give 5-oxo-9-methyl- $\Delta^{1(2),8(9)}$ -octalin-2-(2'-propenoic acid) ethyl ester (0.6 g, yield 35%). This product was subjected to reductive amination in the 15 manner as disclosed in Reference Example 1 to give the titled compound (114 mg, yield 22%).

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.33(\text{d}, 1\text{H}, 15.7\text{Hz}), 6.31(\text{s}, 1\text{H}), 6.86(\text{d}, 1\text{H}, 15.7\text{Hz}), 5.70(\text{t}, 1\text{H}, 3.8\text{Hz}), 4.21(\text{q}, 2\text{H}, 7.1\text{Hz}), 2.83 (\text{dd}, 1\text{H}, 5.6\text{Hz}, 10.4\text{Hz}), 2.2-2.4(\text{m}, 5\text{H}), 2.0-2.18(\text{m}, 1\text{H}), 1.7-1.9 (\text{m}, 2\text{H}), 1.30(\text{t}, 3\text{H}, 7.1\text{Hz}), 0.99(\text{s}, 3\text{H}).$

20 **Reference Example 7**

Preparation of 3-(p-t-butylphenoxy)benzoic acid:

To a solution of 3-bromobenzonitrile (4.35 g, 23.9 mmol) and p-t-butylphenol (4.30 g, 28.7 mmol) in 25 pyridine (45 ml) were added potassium carbonate (6.61 g, 47.8 mmol) and cupric oxide (3.42 g, 43 mmol), and the resultant mixture was refluxed for 46 hours. The reaction mixture was poured into saturated potassium hydrogensulfate solution and extracted with ethyl acetate. The organic layer was concentrated, and the residue was dissolved in ethylene glycol (20 ml), mixed with 2 N potassium hydroxide solution (130 ml) and refluxed for 20 hours. The reaction mixture was poured into chilled water and extracted with diethyl 30 ether under acidic conditions. The organic layer was dried and concentrated. The residue was chromatographed on a silica gel column to give the titled compound (3.46 g, yield 54%).

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.80(\text{d-like}, 1\text{H}, 7.6\text{Hz}), 7.70(\text{t}, 1\text{H}, 1.9\text{Hz}), 7.42(\text{d}, 1\text{H}, 7.6\text{Hz}), 7.36(\text{d}, 2\text{H}, 8.6\text{Hz}), 7.24(\text{dd}, 1\text{H}, 1.9\text{Hz}, 7.6\text{Hz}), 6.94(\text{d}, 2\text{H}, 8.6\text{Hz}), 1.32(\text{s}, 9\text{H}).$

The compounds of the following Reference Examples 8 - 16 were prepared according to Reference 35 Example 7.

Reference Example 8

4-phenoxybenzoic acid:

40 $^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.98(\text{d}, 2\text{H}, 8.8\text{Hz}), 7.34(\text{t}, 2\text{H}, 7.7\text{Hz}), 7.14(\text{t}, 1\text{H}, 7.7\text{Hz}), 7.02(\text{d}, 2\text{H}, 8.8\text{Hz}), 6.93(\text{d}, 2\text{H}, 7.7\text{Hz}), 3.83 (\text{br-s}, 1\text{H}).$

Reference Example 9

45 3-(o-isopropylphenoxy)benzoic acid:

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.77(\text{d}, 1\text{H}, 7.8\text{Hz}), 7.61(\text{br-s}, 1\text{H}), 7.40(\text{d}, 1\text{H}, 8.0\text{Hz}), 7.35(\text{t-like}, 1\text{H}, 4.9\text{Hz}), 7.1-7.2(\text{m}, 3\text{H}), 6.85-6.90(\text{m}, 1\text{H}), 3.15-3.3(\text{m}, 1\text{H}), 1.19, 1.22(\text{each s, each } 3\text{H}).$

Reference Example 10

50 3-(p-isopropylphenoxy)benzoic acid:

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.80(\text{ddd}, 1\text{H}, 1.2\text{Hz}, 1.6\text{Hz}, 7.8\text{Hz}), 7.68(\text{dd}, 1\text{H}, 1.6\text{Hz}, 2.4\text{Hz}), 7.39(\text{dd}, 1\text{H}, 7.8\text{Hz}, 8.0\text{Hz}), 7.18-7.28 (\text{m}, 3\text{H}), 6.90-7.0(\text{m}, 2\text{H}), 2.80-3.00(\text{m}, 1\text{H}), 1.23, 1.26(\text{each s, each } 3\text{H}).$

55 **Reference Example 11**

3-(m-isopropylphenoxy)benzoic acid:

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.83(\text{ddd}, 1\text{H}, 1.2\text{Hz}, 1.5\text{Hz}, 7.7\text{Hz}), 7.72(\text{dd}, 1\text{H}, 1.2\text{Hz}, 2.8\text{Hz}), 7.42(\text{dd},$

EP 0 579 223 A1

1H, 7.9Hz, 8.0Hz), 7.2-7.3(m, 2H), 7.02(dd, -like, 1H, 0.7Hz, 7.2Hz), 6.92(br-s, 1H), 6.82(dd- like, 1.2Hz, 8.2Hz), 2.80-3.00(m, 1H), 1.23, 1.26(each s, each 3H).

Reference Example 12

5 3-[p-(2'-methylpropyl)phenoxy]benzoic acid:

'H-NMR(CDCl₃, 250MHz) δ = 7.89(ddd, 1H, 1.2Hz, 1.7Hz, 7.7Hz), 7.67(dd, 1H, 2.4Hz, 1.7Hz), 7.40(t, 1H, 7.7Hz), 7.21(ddd, 1H, 1.2Hz, 2.4Hz, 7.7Hz), 7.11(d, 2H, 8.5Hz), 6.92(d, 2H, 8.5Hz), 3.75(br-s, 1H), 2.45(d, 2H, 7.2Hz), 1.84(dt, 1H, 6.5Hz, 7.2Hz), 0.90(d, 6H, 6.5Hz).

10

Reference Example 13

3-(p-tolyloxy)benzoic acid:

'H-NMR(CDCl₃, 250MHz) δ = 7.79(ddd, 1H, 1.2Hz, 1.6Hz, 7.7Hz), 7.65(dd, 1H, 1.6Hz, 2.4Hz), 7.39(t, 1H, 7.7Hz), 7.21(ddd, 1.2Hz, 1.8Hz, 7.7Hz), 7.15(d, 2H, 8.6Hz), 6.92(d, 2H, 8.6Hz), 2.34(s, 3H).

Reference Example 14

3-(p-cyclohexylphenoxy)benzoic acid:

'H-NMR(CDCl₃, 250MHz) δ = 7.80(dd-like, 1H, 1.2Hz, 8.0Hz), 7.68(dd, 1H, 1.2Hz, 2.4Hz), 7.39(t, 1H, 8.0Hz), 7.10-7.25(m, 3H), 6.93(d, 2H, 8.6Hz), 2.38-2.58(m, 1H), 1.68-1.96(m, 5H), 1.14-1.52 (m, 5H).

Reference Example 15

25 3-(2',4'-dimethylphenoxy)benzoic acid:

'H-NMR(CDCl₃, 250MHz) δ = 7.81(dd, 1H, 1.6Hz, 7.7Hz), 7.68 (dd, 1H, 1.6Hz, 2.2Hz), 7.40(dd, 1H, 7.7Hz, 7.9Hz), 7.22(dd-like, 2.2Hz, 7.9Hz), 6.77(br-s, 1H), 6.63(br-s, 2H), 2.28(s, 6H).

Reference Example 16

30

3-(5',6',7',8'-tetrahydro-2'-naphthyloxy)benzoic acid:

'H-NMR(CDCl₃, 250MHz) δ = 7.79(br-d, 1H, 7.7Hz), 7.67(br-s, 1H), 7.40(t, 1H, 7.7Hz), 7.23(dd, 1H, 2.4Hz, 7.7Hz), 7.04(d, 1H, 8.0Hz), 6.7-6.8(m, 2H), 2.7-2.8(m, 4H), 1.7-1.9(m, 4H).

35

Reference Example 17

Preparation of 3-bromo-5-(p-isopropylphenoxy)benzoic acid (a) and 3,5-di(p-isopropylphenoxy)benzoic acid (b):

40 To a solution of 3,5-dibromobenzoic acid (3.8 g, 11 mmol) and p-isopropylphenol (3.36 g, 24.2 mmol) in pyridine (46 ml) were added cupric oxide (2.19 g, 27.5 mmol) and potassium carbonate (6.08 g, 44 mmol), and the resultant mixture was refluxed for 5 days. The reaction mixture was allowed to warm up to room temperature, poured into chilled water and extracted with chloroform under acidic conditions. The organic layer was concentrated, and the residue was chromatographed on a silica gel column to give the titled compounds (a) (489 mg, yield 13%) and (b) (1.88 g, yield 44%).

45 (a); 'H-NMR(CDCl₃, 250MHz) δ = 7.19(dd, 1H, each 1.7Hz), 7.60 (dd, 1H, each 1.7Hz), 7.34(dd, 1H, each 1.7Hz), 7.23(d, 2H, 8.2Hz), 6.95(d, 2H, 8.2Hz), 2.8-3.0(m, 1H), 1.24, 1.27(each s, each 3H)
(b); 'H-NMR(CDCl₃, 250MHz) δ = 7.34(d, 2H, 2.1Hz), 7.19(m, 4H, 8.5Hz), 6.94(d, 4H, 8.5Hz), 6.86(d, 1H, 2.1Hz), 2.80-2.95(m, 2H), 1.22, 1.25(each s, each 6H).

50

Reference Example 18

Preparation of 3-dicyclohexylmethoxybenzoic acid:

55 To a solution of 3-hydroxybenzoic acid (1.61 g, 8.2 mmol), dicyclohexylmethanol (1.61 g, 8.2 mmol), triphenylphosphine (2.05 g, 7.82 mmol) and dioxane (35 ml) was added diethyl azodicarboxylate (1.41 ml, 8.94 mmol), and the resultant mixture was refluxed for 3 days. The reaction mixture was allowed to warm up to room temperature, poured into chilled water and extracted with diethyl ether. The organic layer was

EP 0 579 223 A1

concentrated, and the residue was dissolved in a mixture of methanol (20 ml), tetrahydrofuran (6.5 ml) and water (2.0 ml), mixed with potassium hydroxide (1.16 g) and stirred for 2 hours. The mixture was poured into chilled water, acidified, and extracted with ethyl acetate. The organic layer was concentrated, and the resultant residue was recrystallized from hexane to give the titled compound (0.43 g, yield 18 %).

5 ¹H-NMR(CDCl₃, 250MHz) δ = 7.55-7.65(m, 2H), 7.30(dd, 1H, each 8.4Hz), 7.13(ddd, 1H, 0.9Hz, 2.5Hz, 8.4Hz), 3.98(t, 1H, 5.6Hz), 1.50-1.88(m, 12H), 0.96-1.34(m, 10H).

The compounds of the following Reference Examples 19 - 24 were prepared according to Reference Example 18.

10 Reference Example 19

3-(2',4'-dimethyl-3'-pentoxy)benzoic acid:

1H-NMR(CDCl₃, 250MHz) δ = 7.55-7.65(m, 2H), 7.32(t, 1H, 8.2Hz), 7.15(ddd, 1H, 1.0Hz, 2.7Hz, 8.2Hz), 3.94(t, 1H, 5.8Hz), 2.01(dt, 2H, 6.8Hz, 13.3Hz), 0.90-0.98(m, 12H).

15 Reference Example 20

3-(dicyclopropylmethoxy)benzoic acid:

1H-NMR(CDCl₃, 250MHz) δ = 7.50-7.61(m, 2H), 7.28(t, 1H, 8.2Hz), 7.10(ddd, 1H, 1.2Hz, 2.7Hz, 8.2Hz), 20 3.88(s, 3H), 3.49(t, 1H, 6.9Hz), 1.0-1.2(m, 2H), 0.4-0.6(m, 4H), 0.2-0.4(m, 4H).

Reference Example 21

3-(cyclooctyloxy)benzoic acid:

25 1H-NMR(CDCl₃, 250MHz) δ = 7.66(ddd, 1H, 1.2Hz, 2.3Hz and 8.0Hz), 7.57(dd, 1H, 2.3Hz, 1.7Hz), 7.34(t, 8Hz), 7.09(ddd, 1.2Hz, 1.7Hz, 8.0Hz), 4.47(4.0Hz, 7.9Hz, 11.8Hz), 1.4-2.0(m, 14H).

Reference Example 22

3-(4-heptyloxy)benzoic acid:

30 1H-NMR(CDCl₃, 250MHz) δ = 7.64(ddd, 1.2Hz, 2.0Hz, 8.0Hz), 7.59(dd, 1H, 1.2Hz, 2.3Hz), 7.33(t, 1H, 8.0Hz), 7.10(ddd, 1.2Hz, 2.3Hz, 8.9Hz), 4.31(dt, 1H, 5.9Hz, 11.6Hz), 1.3-1.8(m, 8H), 0.91 (t, 6H, 7.0Hz).

Reference Example 23

35 3-(cyclohexylmethoxy)benzoic acid:

1H-NMR(CDCl₃, 250MHz) δ = 7.68(1H, 2.2Hz, 7.7Hz), 7.60(d, 1H, 2.2Hz), 7.35(t, 1H, 7.7Hz), 7.12(dt, 1H, 2.0Hz, 7.7Hz), 3.79(d, 2H, 6.0Hz), 1.6-1.9(m, 6H), 1.0-1.4(m, 5H).

40 Reference Example 24

3-cycloheptyloxy-4-methoxybenzoic acid:

45 1H-NMR(CDCl₃, 250MHz) δ = 7.63(dd, 1H, 2.0Hz, 8.5Hz), 7.50(d, 1H, 2.0Hz), 6.86(d, 1H, 8.5Hz), 4.42(ddd, 1H, 4.3Hz, 8.5Hz and 12.6Hz), 3.88(s, 3H), 3.86(s, 3H), 1.96-2.05(m, 2H), 1.62-1.94(m, 4H), 1.33 -1.62(m, 6H).

Reference Example 25

Preparation of 3-isopropyl-4-cycloheptylbenzoic acid:

50 To a solution of 3,4-dihydroxybenzoic acid ethyl ester (2.0 g, 11 mmol) in acetone (450 l) were added potassium carbonate (2.28 g, 16.5 mmol) and isopropylamide (1.03 ml, 11 mmol), and the resultant mixture was stirred at room temperature for 2 days. The reaction mixture was poured into chilled water and extracted with ethyl acetate under acidic conditions. The organic layer was concentrated, and the residue 55 was chromatographed on a silica gel column to give a mixture (617 mg) of 3-hydroxy-4-isopropoxybenzoic acid ethyl ester and its position isomer, 4-hydroxy-3-isopropoxybenzoic acid ethyl ester (6:1). There was observed NOE of protons at 1'-position of the isopropyl group and 5-position. The reaction was effected using this mixture (617 mg) in the same manner as in Reference Example 18, and the resultant product was

EP 0 579 223 A1

recrystallized from hexane to give the titled compound (370 mg, yield 12%).

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.68(\text{dd}, 1\text{H}, 2.0\text{Hz}, 8.4\text{Hz}), 7.58(\text{d}, 1\text{H}, 2.0\text{Hz}), 6.90(\text{d}, 1\text{H}, 8.4\text{Hz}), 4.55-4.70(\text{m}, 1\text{H}), 4.35-4.50(\text{m}, 1\text{H}), 1.9-2.1(\text{m}, 2\text{H}), 1.3-1.9(\text{m}, 10\text{H}), 1.34-1.37(\text{each s, each } 3\text{H})$.

5 Reference Example 26

Preparation of 3-cyclohexylmethylbenzoic acid:

To a solution of 3-carboxybenzaldehyde (2.65 g, 17.7 mmol) in tetrahydrofuran (2.65 ml) was added a
10 solution of cyclohexyl magnesium bromide in diethyl ether (53 mmol, 30 ml). This mixture was added to 3 N hydrochloric acid (500 ml) and extracted with chloroform. The organic layer was concentrated, and the residue was chromatographed on a silica gel column to give the condensed product (1.35 g). The product was dissolved in methanol (27 ml), mixed with palladium hydroxide (135 mg) and hydrogenated. The reaction mixture was filtered through Celite, and the filtrate was concentrated. The residue was recrystallized from hexane to give the titled compound (0.84 g, yield 22 %).

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.85-8.00(\text{m}, 2\text{H}), 7.30-7.40(\text{m}, 2\text{H}), 2.54(\text{d}, 2\text{H}, 6.9\text{Hz}), 1.5-1.8(\text{m}, 6\text{H}), 1.1-1.3(\text{m}, 3\text{H}), 0.80-1.05(\text{m}, 2\text{H})$.

20 Reference Example 27

Preparation of 3-cycloheptylmethylbenzoic acid:

The titled compound was prepared according to Reference Example 26.

$^1\text{H-NMR}(\text{CDCl}_3, 250\text{MHz})\delta = 7.85-7.95(\text{m}, 2\text{H}), 7.30-7.40(\text{m}, 2\text{H}), 2.56(\text{d}, 2\text{H}, 7.2\text{Hz}), 1.1-1.9(\text{m}, 13\text{H})$.

25 **EXPERIMENT**

Human steroid 5α -reductase inhibiting effect was examined on the compounds of the present invention using human prostatic tissue.

30 Frozen human prostate was thawed on ice and minced into small pieces (~ 5mm³). The minced tissue was homogenized with a Polytron homogenizer (Kinematica, Switzerland) in 40 vol. of 0.1 M phosphate buffer at pH 5.5. The suspension was used as the source of 5α -reductase.

The reaction tube contained 50nM ($1\alpha,2\alpha$ - $^3\text{H}(N)$)tetstosterone (2.04 TBq/mmol, New England Nuclear, MA), a predetermined amount of an inhibitor, 0.5 mg of NADPH and 0.5 ml of prostatic suspension in a total 35 final volume of 1 ml. For this test, DMSO solutions containing various amount of the inhibitor were employed.

40 Each reaction solution was incubated at 37 °C for 10 minutes and ethyl acetate (2 ml) chilled with ice was added for stopping the reaction. The organic layer was transferred to another test tube and evaporated to dryness under nitrogen. Each 50 μg of non radioactive testosterone, DHT, estradiol, androstenedione and androstandiol was added to the residue, and the wall of each test tube was washed with ethyl acetate (0.1 ml). The content of the tube was evaporated again and the residue was dissolved in chloroform (50 μl) and then applied to Whatman LK60DF silica plates (Whatman, NJ), developing in chloroform:methanol (50:1). The bands of the substrates and products were identified by fluorescence (254 nm) and iodine, and the relevant silica gel was collected by scraping, mixed with toluene based liquid scinilator, and the radioactivity was counted. The radioactivities of DHT and androstandiol were combined for the calculation of 5α -reductase activity. The conversion rate from testosterone to DHT and androstandiol was 35%-45% in the control group which only the solvent was added in place of inhibitors. The radioactivity for the control group was postulated as 100%.

45 IC_{50} were calculated from the data obtained from the above experiments. Table 5 shows the test results.

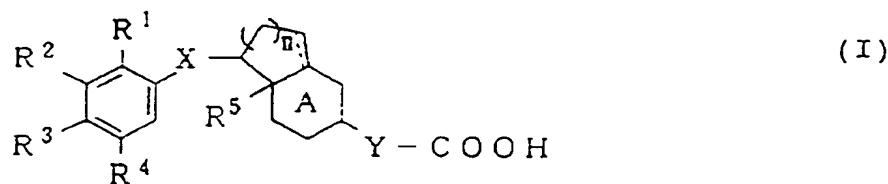
Table 5

	Experiment No.	IC ₅₀ (μ M)	Experiment No.	IC ₅₀ (μ l)
5	1	3.3	23	0.30
	2	0.83	31	0.41
	4	0.81	35	0.16
	5	0.46	37	0.30
	7	0.21	39	0.39
	9	0.52	42	0.77
	11	0.25	43	0.23
	13	0.66	45	0.61
	14	0.44	46	0.30
	15	4.21	47	0.16
15	17	0.18	48	0.065
	20	0.27	51	0.91
	22	0.52	54	0.16

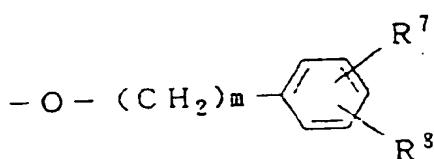
20 Table 5 shows that the compounds of the present invention and salts thereof are useful as 5 α -reductase inhibitors. Accordingly, they would be useful as therapeutic agents to diseases such as benign prostatic hyperplasia, acne, seborrhea, female hirsutism, prostatic cancer, male alopecia or the like, in which the reduction of DHT activity is expected to be effective for therapeutic treatment thereof.

25 **Claims**

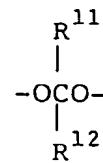
1. Carboxylic acid derivatives of the following general formula (I):



Wherein R¹, R², R³ and R⁴ each independently represent hydrogen atom, halogen atom, adamantyl group, optionally substituted C₁ - C₁₄ alkyl group, optionally substituted C₃ - C₁₀ cycloalkyl group, 40 optionally substituted C₁ - C₁₄ alkoxy group, optionally substituted heterocyclic group, - OR⁶ (R⁶ represents hydrogen atom, adamantyl group, optionally substituted C₃ - C₁₀ cycloalkyl group or 45 optionally substituted heterocyclic group), or a group of the formula:



wherein R⁷ and R⁸ each independently represent hydrogen atom, C₁ - C₆ alkyl group, C₃ - C₈ cycloalkyl group, - CONR⁹N¹⁰ (R⁹ and R¹⁰ each independently represent hydrogen atom or C₁ - C₆ alkyl group) or, when R⁷ and R⁸ are adjacent, they may form C₁ - C₆ alkylene group, and m represents 55 0 or 1, or, the adjacent two substituents selected from R¹, R², R³ and R⁴ may form a group of the formula:



wherein R¹¹ and R¹² each independently represent hydrogen atom, C₁ - C₆ alkyl group or C₃ - C₈ cycloalkyl group, or they may form, taken together, C₂ - C₈ alkylene group, a group of the formula -OCH₂CH₂O- or optionally substituted C₃ - C₄ alkylene group,

R^5 represents hydrogen atom or $C_1 - C_5$ alkyl group.

X represents -CONR¹³- or -SO₂NR¹³- (R¹³ represents hydrogen atom or C₁ - C₆ alkyl group).

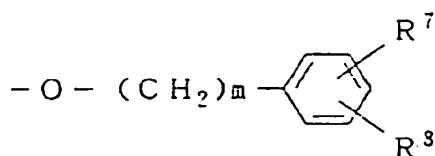
Y represents a single bond, $-\text{OCH}_2-$ or $-\text{CH}=\text{CH}-$.

A ring may form a benzene ring, cyclohexene ring or cyclohexadiene ring.

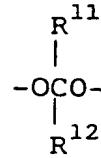
the dotted line represents a single bond or double bond, and

n represents 1 or 2, with the proviso that when the carbon atom to which R⁵ is attached has a double bond, then R⁵ is not present, or pharmaceutically acceptable salts thereof.

20 2. A compound according to Claim 1, in which R¹, R², R³ and R⁴ each independently represent hydrogen atom; halogen atom; adamantyl group; C₁ - C₁₂ alkyl group optionally having one or more substituents selected from 5 or 6 membered heterocyclic group containing 1 or 2 hetero atoms selected from oxygen atom, sulfur atom and nitrogen atom, C₃ - C₁₀ cycloalkyl group and adamantyl group; C₃ - C₁₀ cycloalkyl group optionally having one or more C₁ - C₆ alkyl groups; C₁ - C₁₄ alkoxy group optionally having two or more phenyl groups or one or more substituents selected from 5 or 6 membered heterocyclic group containing 1 or 2 hetero atoms selected from oxygen atom, sulfur atom and nitrogen atom, C₃ - C₁₀ cycloalkyl group and adamantyl group; 5 or 6 membered heterocyclic group containing 1 or 2 hetero atoms selected from oxygen atom, sulfur atom and nitrogen atom and optionally having one or more C₁ - C₅ alkyl groups; -OR⁶ [R⁶ represents hydrogen atom; adamantyl group; C₃ - C₁₀ cycloalkyl group optionally having one or more C₁ - C₆ alkyl groups; or 5 or 6 membered heterocyclic group containing 1 or 2 hetero atoms selected from oxygen atom, sulfur atom and nitrogen atom, and optionally having one or more substituents selected from C₁ - C₆ alkyl group and C₂ - C₆ acyl group]; or a group of the formula:



45 wherein R⁷ and R⁸ each independently represent hydrogen atom, C₃ - C₈ cycloalkyl group, -CONR⁹R¹⁰ (R⁹ and R¹⁰ each independently represent hydrogen atom or C₁ - C₆ alkyl group), or, when R⁷ and R⁸ are adjacent, they may form C₁ - C₆ alkylene group, and m represents 0 or 1, or, the adjacent two substituents selected from R¹, R², R³ and R⁴ may form a group of the formula:



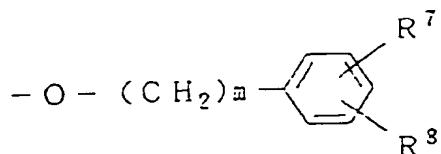
55

wherein R¹¹ and R¹² each independently represent hydrogen atom, C₁ - C₆ alkyl group or C₃ - C₈ cycloalkyl group, or they may form, taken together, C₂ - C₈ alkylene group, a group of the formula:

EP 0 579 223 A1

-OCH₂CH₂O-; or C₃ - C₄ alkylene group optionally having one or more substituents selected from C₁ - C₆ alkyl group and C₅ - C₈ cycloalkyl group.

3. A compound according to Claim 2, in which R¹, R², R³ and R⁴ each independently represent hydrogen atom; halogen atom; C₁ - C₁₄ alkyl group optionally having one or more substituents selected from 5 or 6 membered heterocyclic group containing 1 or 2 hetero atoms selected from oxygen atom, sulfur atom and nitrogen atom, and C₃ - C₆ cycloalkyl group; C₁ - C₁₄ alkoxy group optionally having two or more phenyl groups or one or more C₃ - C₁₀ cycloalkyl groups; -OR⁵ [R⁶ represents C₃ - C₁₀ cycloalkyl group optionally having one or more C₁ - C₆ alkyl groups; or 5 or 6 membered heterocyclic group containing 1 or 2 hetero atoms selected from oxygen atom, sulfur atom and nitrogen atom, and optionally having one or more C₁ - C₆ alkyl group, or a group of the formula:



20 wherein R⁷ and R⁸ each independently represent hydrogen atom, C₃ - C₈ cycloalkyl group, -CONR⁹N¹⁰ (R⁹ and R¹⁰ each independently represents hydrogen atom or C₁ - C₆ alkyl group), or, when R⁷ and R⁸ are adjacent, they may form C₁ - C₆ alkylene group, and m represents 0 or 1, R⁵ represents C₁ - C₆ alkyl group, X represents -CONR¹³- or -SO₂NR¹³- (R¹³ represents hydrogen or C₁ - C₆ alkyl group), and A ring may form a benzene ring or cyclohexene ring.

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4. A compound according to Claim 3, in which R¹, R², R³ and R⁴ each independently represent hydrogen atom; halogen atom; C₁ - C₁₄ alkyl group optionally having one or more C₃ - C₁₀ cycloalkyl group; C₁ - C₁₄ alkoxy group optionally having one or more C₃ - C₁₀ alkyl groups; or -OR⁶ (R⁶ represents C₁ - C₆ cycloalkyl group optionally having one or more C₁ - C₆ alkyl groups), X represents -CONR¹³- (R¹³ represents hydrogen atom), Y represents a single bond, A ring represents a cyclohexene ring, and the dotted line represents a double bond.

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5. A compound according to Claim 4, in which R¹, R², R³ and R⁴ each independently represent hydrogen atom or C₁ - C₁₄ alkoxy group and n represents 2.

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6. A pharmaceutical composition comprising a carboxylic acid derivative according to Claim 1 or a pharmaceutically acceptable salt thereof together with a pharmaceutically acceptable carrier therefor.

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7. A therapeutic drug for treating androgen dependent diseases comprising a carboxylic acid derivative according to Claim 1 or a pharmaceutically acceptable salt thereof as an effective ingredient together with a pharmaceutically acceptable carrier therefor.

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8. A therapeutic drug according to Claim 7, in which the androgen dependent disease is benign prostatic hyperplasia.

9. A therapeutic drug according to Claim 7, in which the androgen dependent disease is acne.

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10. A therapeutic drug according to Claim 7, in which the androgen dependent disease is seborrhea.

11. A therapeutic drug according to Claim 7, in which the androgen dependent disease is female hirsutism.

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12. A therapeutic drug according to Claim 7, in which the androgen dependent disease is male alopecia.



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EUROPEAN SEARCH REPORT

Application Number

EP 93 11 1367 4

DOCUMENTS CONSIDERED TO BE RELEVANT			CLASSIFICATION OF THE APPLICATION (Int.Cl.S)
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	C07C235/54 A61K31/195 C07D211/44 C07D309/12 A61K31/445 A61K31/35 C07C235/62
A	EP-A-0 291 245 (ONO PHARMACEUTICAL CO.) * claims; examples *	1-12	C07C235/54 A61K31/195 C07D211/44 C07D309/12 A61K31/445 A61K31/35 C07C235/62
D	& JP-A-1 156 950 (...) ----		
A	EP-A-0 294 035 (ONO PHARMACEUTICAL CO.) * claims; examples *	1-12	
D	& JP-A-1 139 558 (...) ----		
A	CHEMICAL ABSTRACTS, vol. 103, no. 25, 13 December 1985, Columbus, Ohio, US; abstract no. 215005m, 'Carboxamide derivatives' page 865 ; * abstract * & JP-A-60 097 946 (ONO PHARMACEUTICAL CO.) -----	1-12	
The present search report has been drawn up for all claims			TECHNICAL FIELDS SEARCHED (Int.Cl.S)
			C07C
Place of search			Examiner
THE HAGUE	Date of completion of the search	29 November 1993	Seufert, G
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone	T : theory or principle underlying the invention		
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